

L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2003 ACS
 AB Z(NHZ1NHR)2 [I; R = H or alk(en)yl; Z = BAB; A,Z1 = (cyclo)alk(en)ylene,
 arylene; B = bond or alk(en)ylene] were prepd. Thus, N,N'-
 bis(mesitylsulfonyl)-cis-1,2-cyclobutanediamine (prepn. given) was
 N-alkylated by Br(CH2)3NEtSO2C6H2Me3-2,4,6 to give, after deprotection,
 Z[NH(CH2)3NHET]2 (Z = cis-1,2-cyclobutylene). Data for biol. activity of
 I were given in graphic form.

ACCESSION NUMBER: 1999:212803 CAPLUS
 DOCUMENT NUMBER: 130:252086
 TITLE: Preparation of conformationally restricted spermine
 analogs as antineoplastic agents
 INVENTOR(S): Frydman, Benjamin J.; Marton, Laurence J.; Reddy,
 Vendohar K.; Valasinas, Aldonia L.; Witiak, Donald T.
 PATENT ASSIGNEE(S): Wisconsin Alumni Research Foundation, USA
 SOURCE: U.S., 41 pp.
 CODEN: USXXAM
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

ALL CMPOS

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5889061	A	19990330	US 1997-951015	19971015 <--
US 6392098	B1	20020521	US 1999-280278	19990329
PRIORITY APPLN. INFO.: MARPAT 130:252086			US 1997-951015	A1 19971015
OTHER SOURCE(S):				
REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT				

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 E1 THROUGH E69 ASSIGNED

	SINCE FILE	TOTAL
	ENTRY	SESSION
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FULL ESTIMATED COST		
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STRUCTURE FILE UPDATES: 13 JUL 2003 HIGHEST RN 547695-13-6
 DICTIONARY FILE UPDATES: 13 JUL 2003 HIGHEST RN 547695-13-6

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2003

Please note that search-term pricing does apply when
 conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

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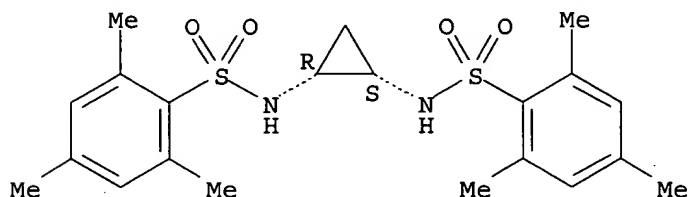
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=> d scan

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-(1R,2S)-1,2-cyclopropanediylbis[2,4,6-trimethyl-,
 rel- (9CI)
 MF C21 H28 N2 O4 S2

Relative stereochemistry.

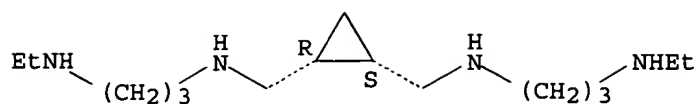


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):68

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedimethanamine, N,N'-bis[3-(ethylamino)propyl]-,
 tetrahydrochloride, (1R,2S)-rel- (9CI)
 MF C15 H34 N4 . 4 Cl H

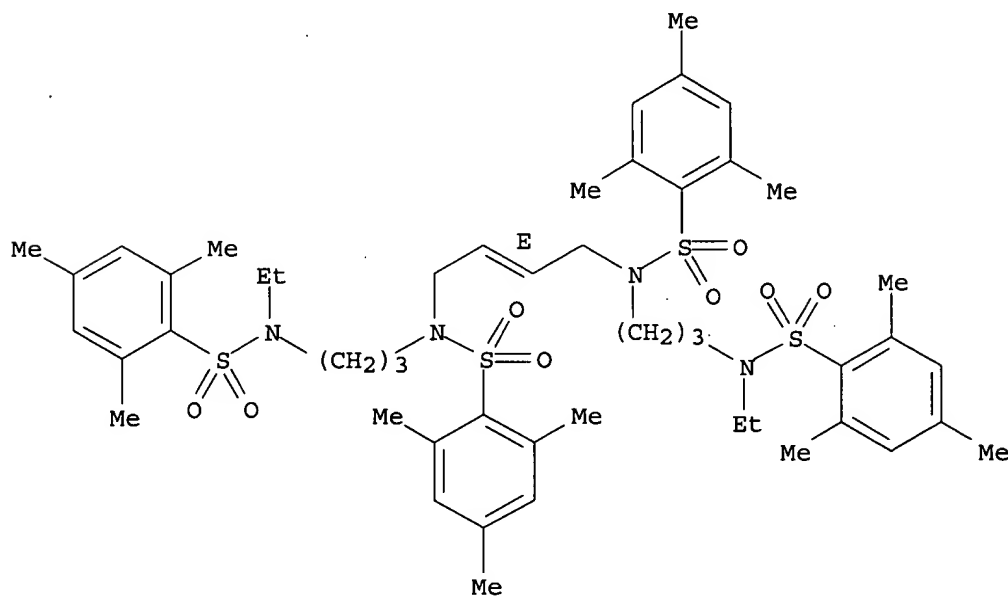
Relative stereochemistry.



●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-(2E)-2-butene-1,4-diylbis[N-[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl- (9CI)
 MF C50 H72 N4 O8 S4

Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

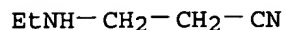
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 2-Propenenitrile (9CI)
 MF C3 H3 N
 CI COM



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS

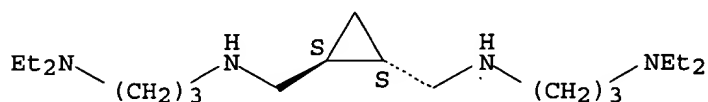
IN Propanenitrile, 3-(ethylamino)- (9CI)
MF C5 H10 N2
CI COM



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclopropanedimethanamine, N,N'-bis[3-(diethylamino)propyl]-,
tetrahydrochloride, (1R,2R)-rel- (9CI)
MF C19 H42 N4 . 4 Cl H

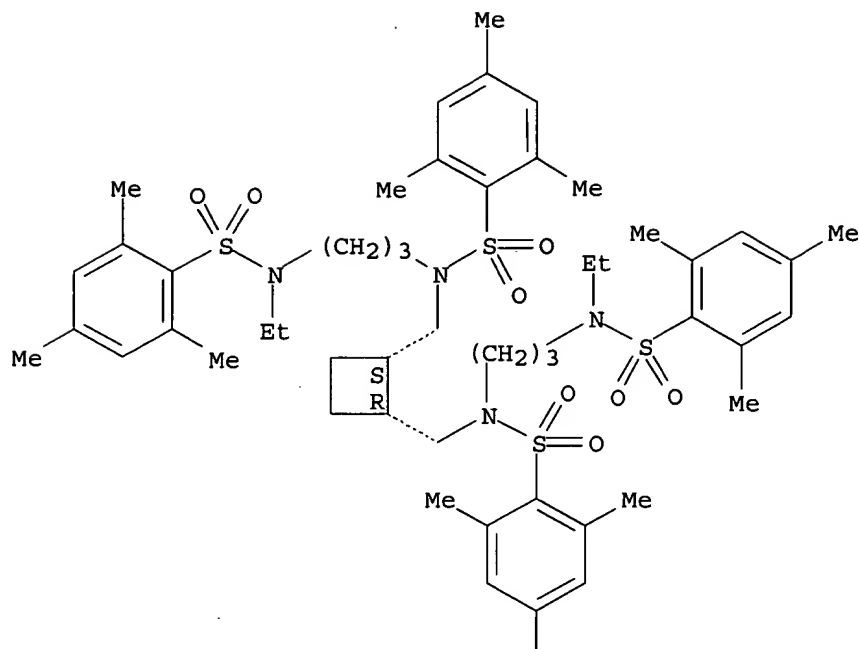
Relative stereochemistry.



●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN Benzenesulfonamide, N,N'-[(1R,2S)-1,2-cyclobutanediylbis(methylene)]bis[N-
[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl-,
rel- (9CI)
MF C52 H76 N4 O8 S4

Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedimethanaminium, N,N,N,N',N',N'-hexamethyl-, dichloride,
 (1R,2S)-rel- (9CI)
 MF C11 H26 N2 . 2 Cl

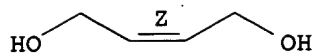
Relative stereochemistry.



●2 Cl⁻

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 2-Butene-1,4-diol, (2Z)- (9CI)
 MF C4 H8 O2
 CI COM

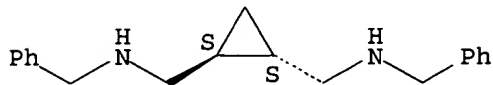
Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedimethanamine, N,N'-bis(phenylmethyl)-, (1R,2R)-rel- (9CI)
 MF C19 H24 N2

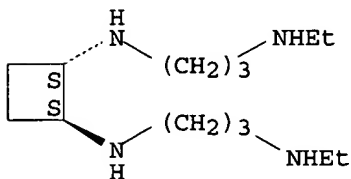
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanediamine, N,N'-bis[3-(ethylamino)propyl]-, tetrahydrochloride, (1R,2R)-rel- (9CI)
 MF C14 H32 N4 . 4 Cl H

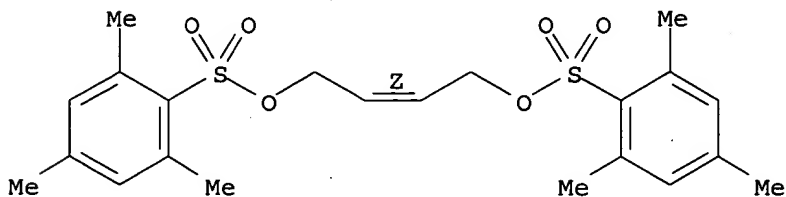
Relative stereochemistry.



● 4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonic acid, 2,4,6-trimethyl-, (2Z)-2-butene-1,4-diyl ester (9CI)
 MF C22 H28 O6 S2

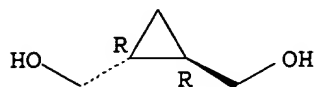
Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclopropanedimethanol, (1R,2R)-rel- (9CI)
MF C5 H10 O2

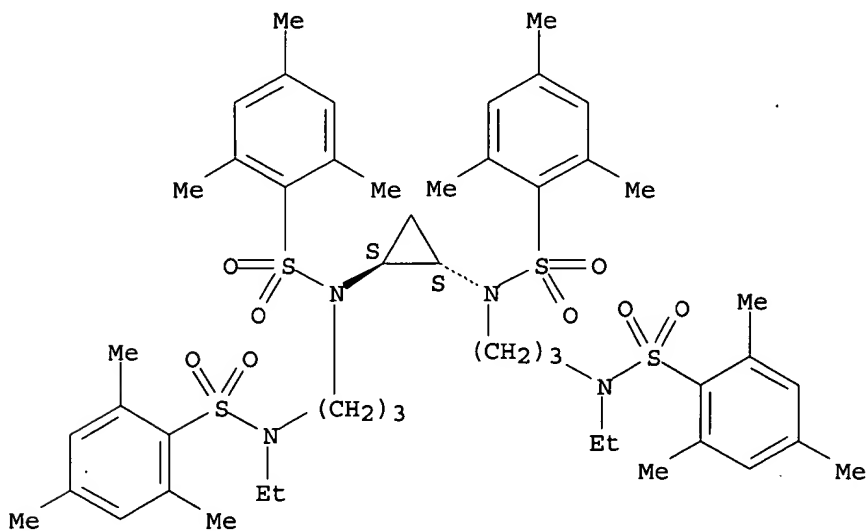
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN Benzenesulfonamide, N,N'-(1R,2R)-1,2-cyclopropanediylbis[N-[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl-, rel- (9CI)
MF C49 H70 N4 O8 S4

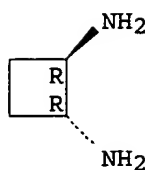
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclobutanediamine, dihydrochloride, (1R,2R)-rel- (9CI)
MF C4 H10 N2 . 2 Cl H

Relative stereochemistry.

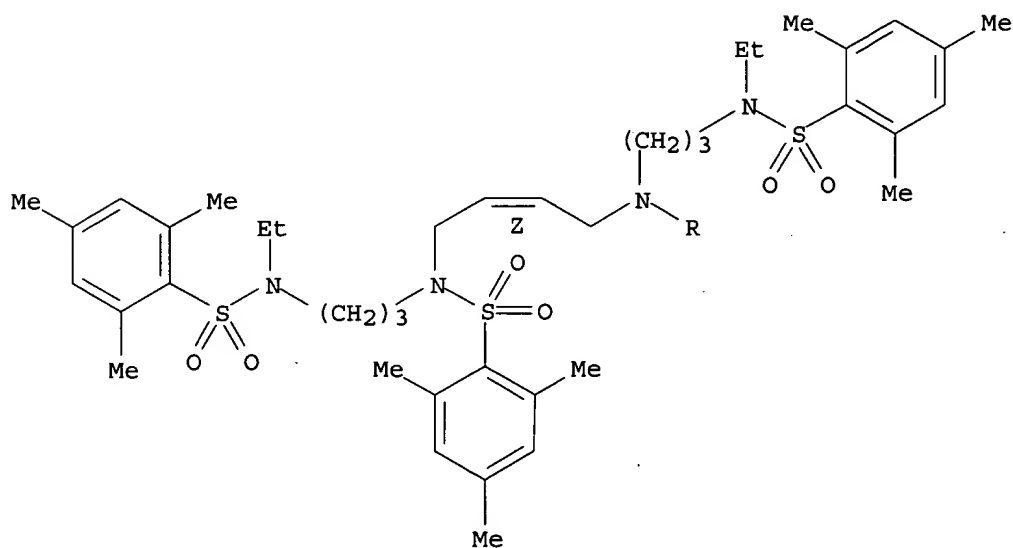


● 2 HCl

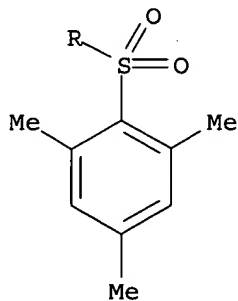
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-(2Z)-2-butene-1,4-diylbis[N-[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl- (9CI)
 MF C50 H72 N4 O8 S4

Double bond geometry as shown.

PAGE 1-A



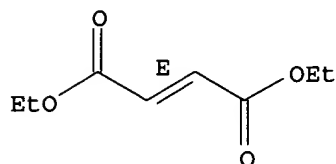
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PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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MF C8 H12 O4
CI COM

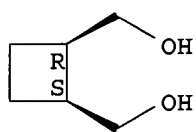
Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclobutanedimethanol, (1R,2S)-rel- (9CI)
MF C6 H12 O2

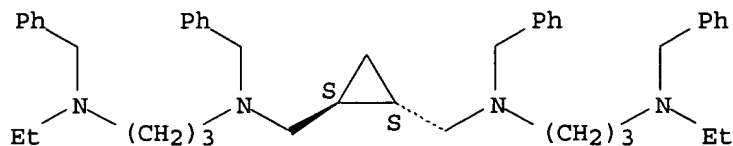
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclopropanedimethanamine, N,N'-bis[3-[ethyl(phenylmethyl)amino]propyl]-N,N'-bis(phenylmethyl)-, (1R,2R)-rel- (9CI)
MF C43 H58 N4

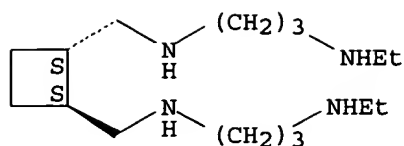
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclobutanedimethanamine, N,N'-bis[3-(ethylamino)propyl]-, tetrahydrochloride, (1R,2R)-rel- (9CI)
MF C16 H36 N4 . 4 Cl H

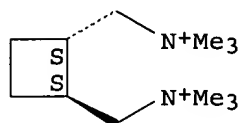
Relative stereochemistry.



●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanedimethanaminium, N,N,N,N',N',N'-hexamethyl-, dichloride,
 (1R,2R)-rel- (9CI)
 MF C12 H28 N2 . 2 Cl

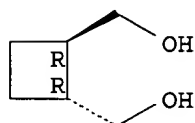
Relative stereochemistry.



●2 Cl⁻

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanedimethanol, (1R,2R)-rel- (9CI)
 MF C6 H12 O2

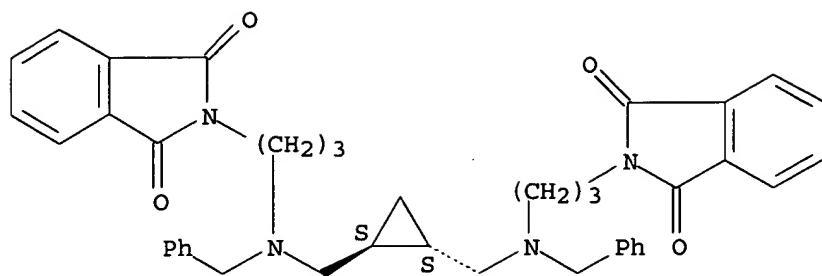
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1H-Isoindole-1,3(2H)-dione, 2,2'-[(1R,2R)-1,2-cyclopropanediylbis[methylene[(phenylmethyl)imino]-3,1-propanediyl]]bis-,
 rel- (9CI)
 MF C41 H42 N4 O4

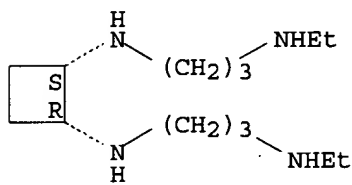
Relative stereochemistry.



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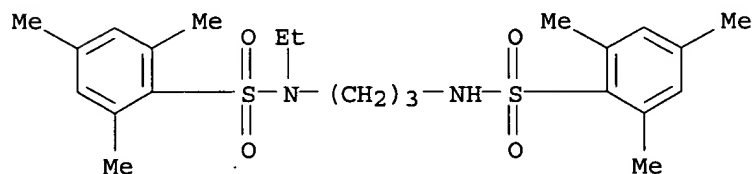
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanediylbis[3-(ethylamino)propyl]-,
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 MF C14 H32 N4 . 4 Cl H

Relative stereochemistry.



● 4 HCl

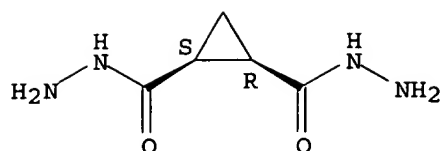
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N-ethyl-2,4,6-trimethyl-N-[3-[[[(2,4,6-
 trimethylphenyl)sulfonyl]amino]propyl]- (9CI)
 MF C23 H34 N2 O4 S2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedicarboxylic acid, dihydrazide, (1R,2S)-rel- (9CI)
 MF C5 H10 N4 O2

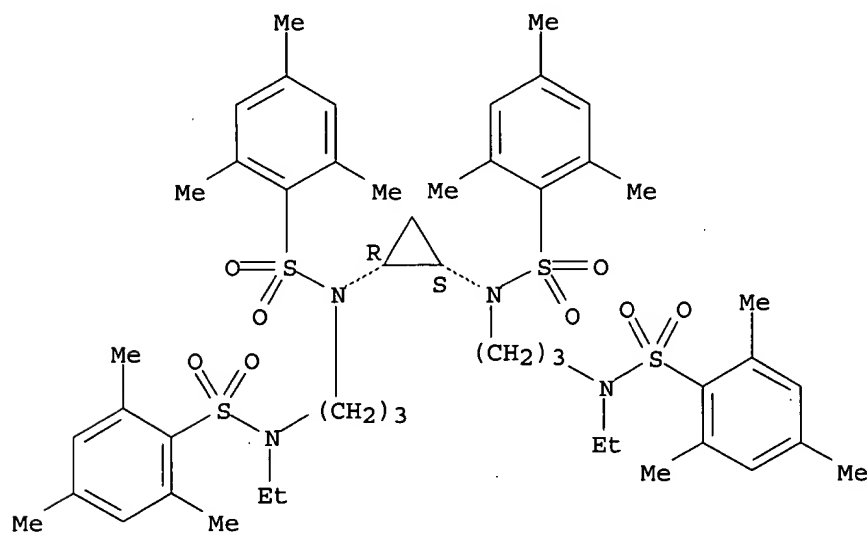
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-(1R,2S)-1,2-cyclopropanediylbis[N-[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl-,
 rel- (9CI)
 MF C49 H70 N4 O8 S4

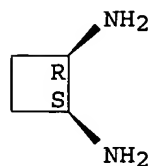
Relative stereochemistry.



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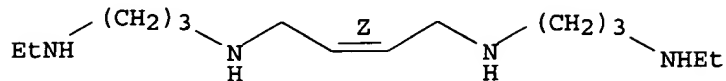
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanediylbis, dihydrochloride, (1R,2S)-rel- (9CI)
 MF C4 H10 N2 . 2 Cl H

Relative stereochemistry.



L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 2-Butene-1,4-diamine, N,N'-bis[3-(ethylamino)propyl]-, tetrahydrochloride,
 (2Z) - (9CI)
 MF C14 H32 N4 . 4 Cl H

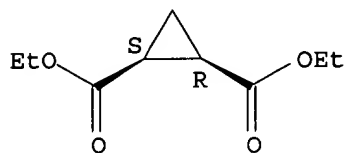
Double bond geometry as shown.



●4 HCl

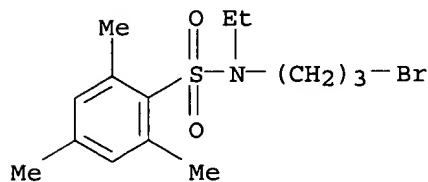
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedicarboxylic acid, diethyl ester, (1R,2S)-rel- (9CI)
 MF C9 H14 O4

Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

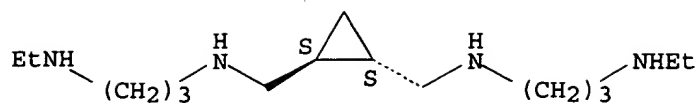
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N-(3-bromopropyl)-N-ethyl-2,4,6-trimethyl- (9CI)
 MF C14 H22 Br N O2 S



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedimethanamine, N,N'-bis[3-(ethylamino)propyl]-,
 tetrahydrochloride, (1R,2R)-rel- (9CI)
 MF C15 H34 N4 . 4 Cl H

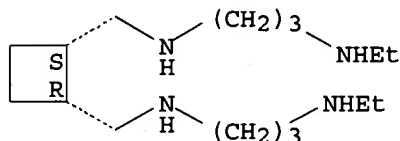
Relative stereochemistry.



●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanedimethanamine, N,N'-bis[3-(ethylamino)propyl]-,
 tetrahydrochloride, (1R,2S)-rel- (9CI)
 MF C16 H36 N4 . 4 Cl H

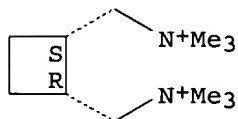
Relative stereochemistry.



●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanedimethanaminium, N,N,N,N',N',N'-hexamethyl-, dichloride,
 (1R,2S)-rel- (9CI)
 MF C12 H28 N2 . 2 Cl

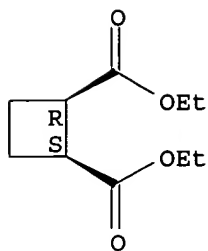
Relative stereochemistry.



●2 Cl⁻

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanedicarboxylic acid, diethyl ester, (1R,2S)-rel- (9CI)
 MF C10 H16 O4

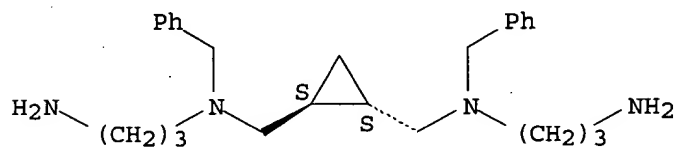
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedimethanamine, N,N'-bis(3-aminopropyl)-N,N'-bis(phenylmethyl)-, (1R,2R)-rel- (9CI)
 MF C25 H38 N4

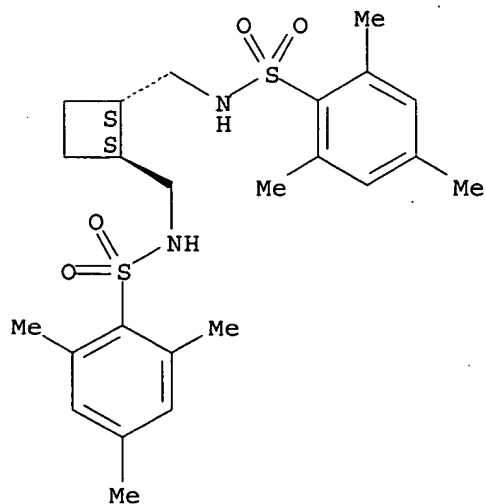
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-[(1R,2R)-1,2-cyclobutanediylbis(methylene)]bis[2,4,6-trimethyl-, rel- (9CI)
 MF C24 H34 N2 O4 S2

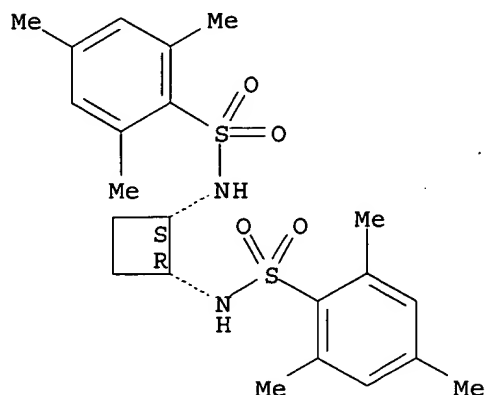
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN Benzenesulfonamide, N,N'-(1R,2S)-1,2-cyclobutanediylbis[2,4,6-trimethyl-,
rel- (9CI)
MF C22 H30 N2 O4 S2

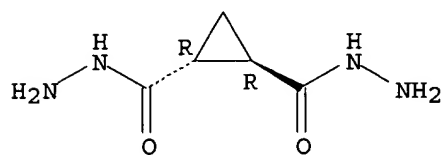
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclopropanedicarboxylic acid, dihydrazide, (1R,2R)-rel- (9CI)
MF C5 H10 N4 O2
CI COM

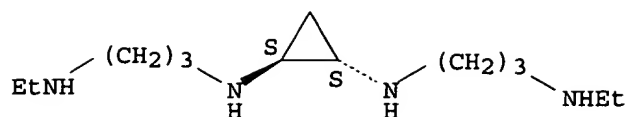
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclopropanediamine, N,N'-bis[3-(ethylamino)propyl]-,
tetrahydrochloride, (1R,2R)-rel- (9CI)
MF C13 H30 N4 . 4 Cl H

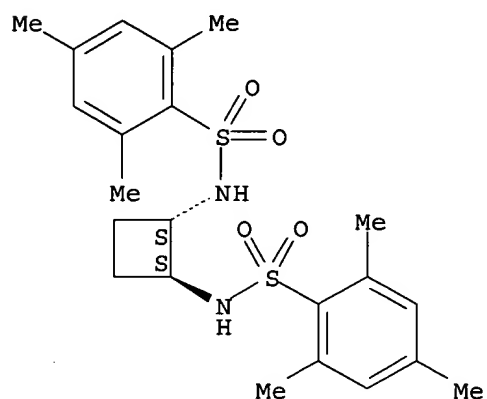
Relative stereochemistry.



●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-(1R,2R)-1,2-cyclobutanediylbis(2,4,6-trimethyl-,
 rel- (9CI)
 MF C22 H30 N2 O4 S2

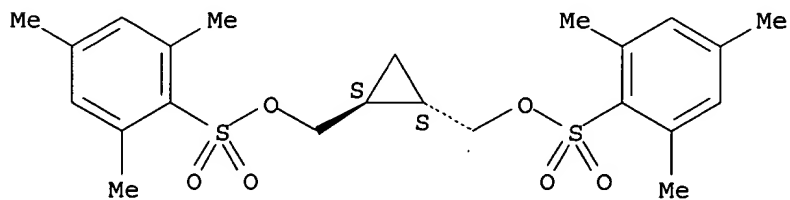
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

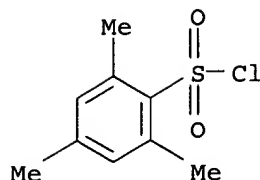
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonic acid, 2,4,6-trimethyl-, (1R,2R)-1,2-
 cyclopropanediylbis(methylene) ester, rel- (9CI)
 MF C23 H30 O6 S2

Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

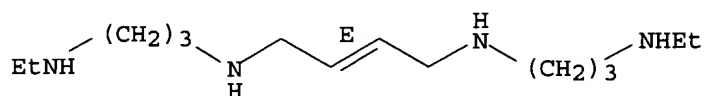
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonyl chloride, 2,4,6-trimethyl- (9CI)
 MF C9 H11 Cl O2 S



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 2-Butene-1,4-diamine, N,N'-bis[3-(ethylamino)propyl]-, tetrahydrochloride,
 (2E)- (9CI)
 MF C14 H32 N4 . 4 Cl H

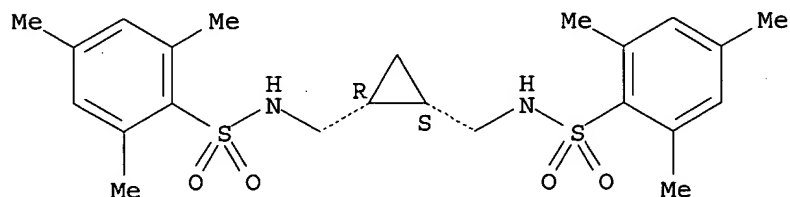
Double bond geometry as shown.



● 4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-[(1R,2S)-1,2-cyclopropanediylbis(methylene)]bis[2,
 4,6-trimethyl-, rel- (9CI)
 MF C23 H32 N2 O4 S2

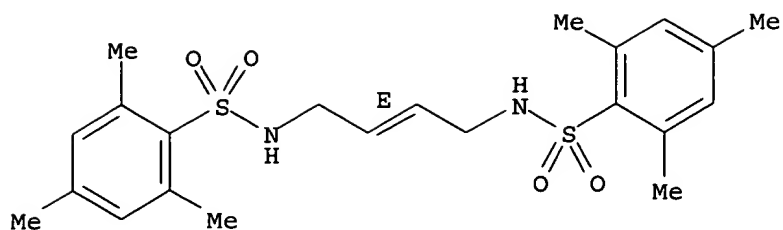
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-(2E)-2-butene-1,4-diylbis[2,4,6-trimethyl- (9CI)
 MF C22 H30 N2 O4 S2

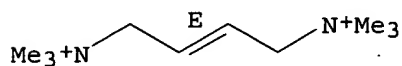
Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 2-Butene-1,4-diaminium, N,N,N,N',N',N'-hexamethyl-, dichloride, (2E)-
 (9CI)
 MF C10 H24 N2 . 2 Cl

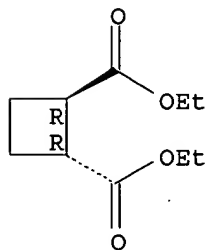
Double bond geometry as shown.



● 2 Cl⁻

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclobutanedicarboxylic acid, diethyl ester, (1R,2R)-rel- (9CI)
 MF C10 H16 O4

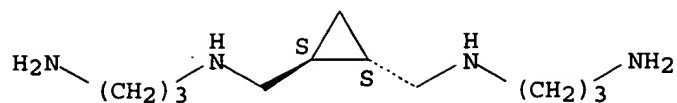
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedimethanamine, N,N'-bis(3-aminopropyl)-,
 tetrahydrochloride, (1R,2R)-rel- (9CI)
 MF C11 H26 N4 . 4 Cl H

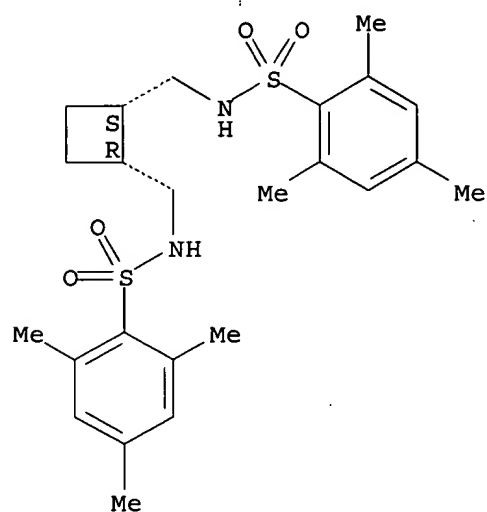
Relative stereochemistry.



●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-[(1R,2S)-1,2-cyclobutanediylbis(methylene)]bis[2,4,6-trimethyl-, rel- (9CI)
 MF C24 H34 N2 O4 S2

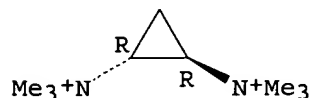
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanediaminium, N,N,N,N',N',N'-hexamethyl-, dichloride, (1R,2R)-rel- (9CI)
 MF C9 H22 N2 . 2 Cl

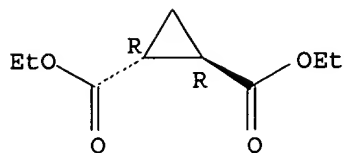
Relative stereochemistry.



●2 Cl⁻

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedicarboxylic acid, diethyl ester, (1R,2R)-rel- (9CI)
 MF C9 H14 O4

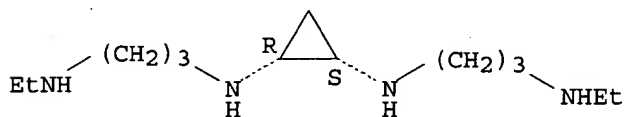
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclopropanediamine, N,N'-bis[3-(ethylamino)propyl]-,
tetrahydrochloride, (1R,2S)-rel- (9CI)
MF C13 H30 N4 . 4 Cl H

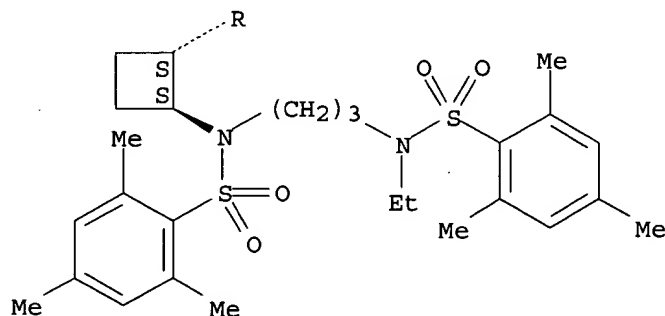
Relative stereochemistry.



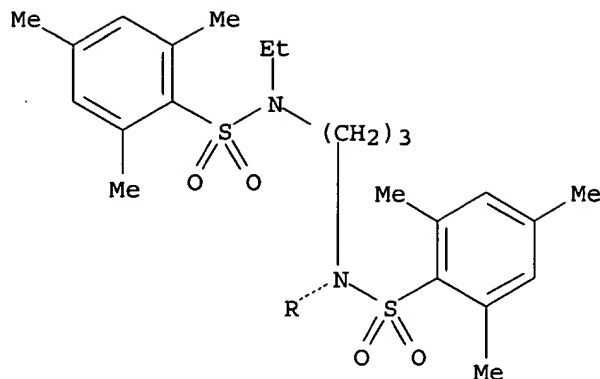
●4 HCl

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN Benzenesulfonamide, N,N'-(1R,2R)-1,2-cyclobutanediylbis[N-[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl-, rel- (9CI)
MF C50 H72 N4 O8 S4

Relative stereochemistry.



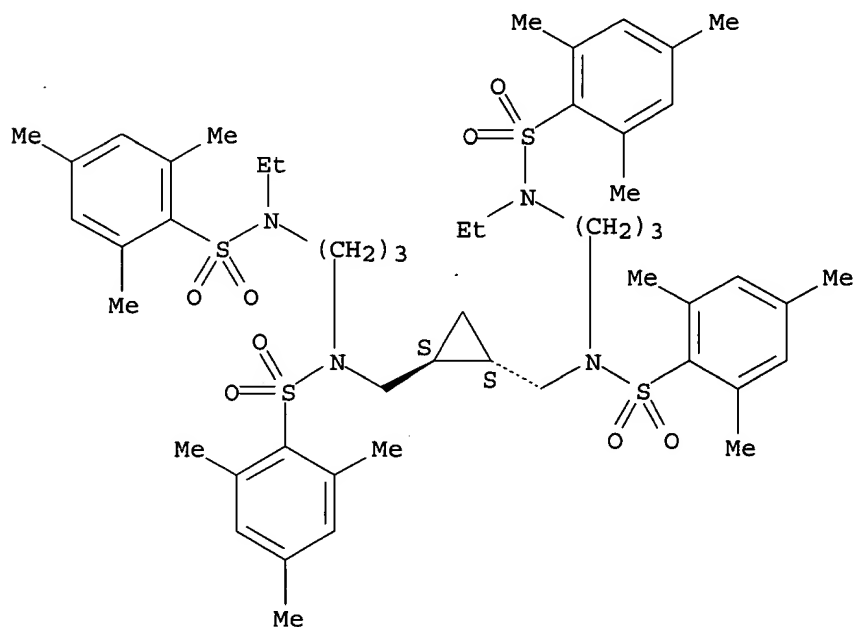
PAGE 1-A



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-[1,2-cyclopropanediylbis(methylene)]bis[N-[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl-, (1R,2R)-rel- (9CI)
 MF C51 H74 N4 O8 S4

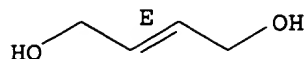
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 2-Butene-1,4-diol, (2E)- (9CI)
 MF C4 H8 O2
 CI COM

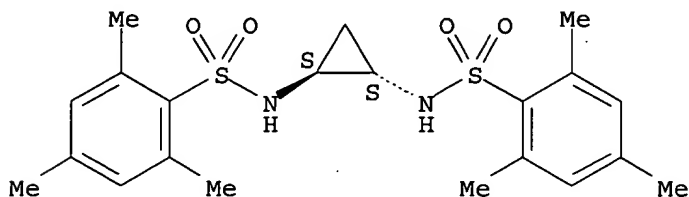
Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN Benzenesulfonamide, N,N'-(1R,2R)-1,2-cyclopropanediylbis[2,4,6-trimethyl-,
rel- (9CI)
MF C21 H28 N2 O4 S2

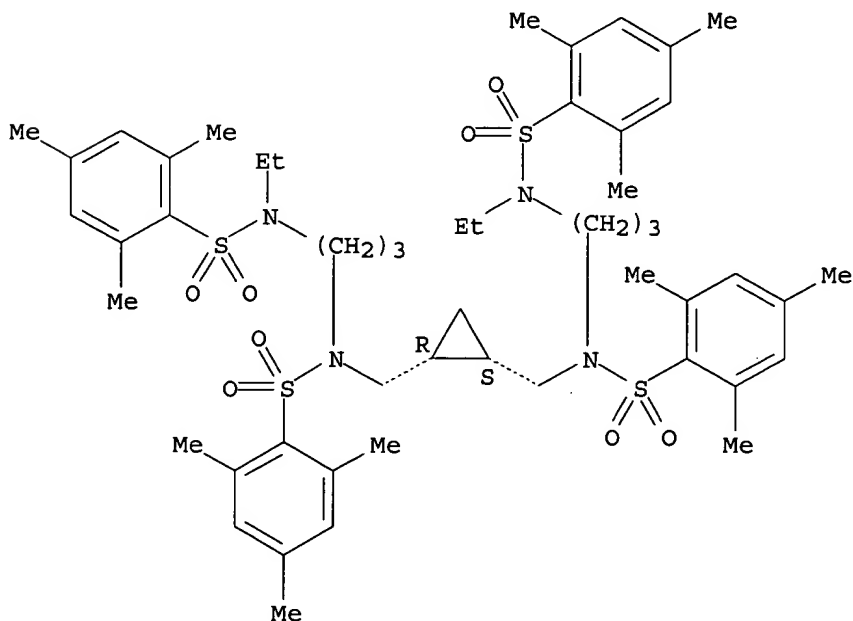
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN Benzenesulfonamide, N,N'-[(1R,2S)-1,2-cyclopropanediylbis(methylene)]bis[N-
[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl-,
rel- (9CI)
MF C51 H74 N4 O8 S4

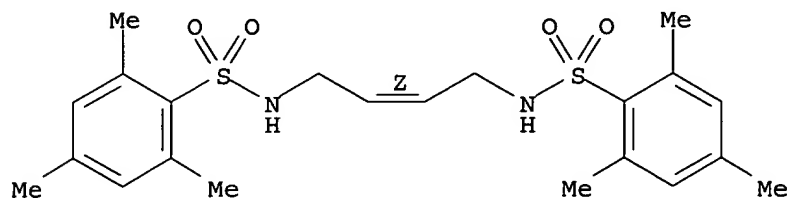
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN Benzenesulfonamide, N,N'-(2Z)-2-butene-1,4-diylbis[2,4,6-trimethyl- (9CI)
MF C22 H30 N2 O4 S2

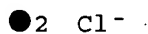
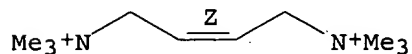
Double bond geometry as shown.



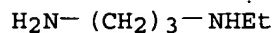
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 2-Butene-1,4-diaminium, N,N,N,N',N',N'-hexamethyl-, dichloride, (2Z)- (9CI)
MF C10 H24 N2 . 2 Cl

Double bond geometry as shown.



L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,3-Propanediamine, N-ethyl- (7CI, 8CI, 9CI)
MF C5 H14 N2
CI COM



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN 1,2-Cyclopropanedimethanamine, N,N'-bis[3-(diethylamino)propyl]-N,N'-bis(phenylmethyl)-, (1R,2R)-rel- (9CI)
MF C33 H54 N4

Relative stereochemistry.



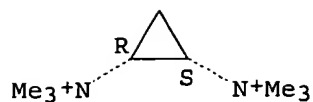
Relative stereochemistry.

The chemical structure shows a central 1,4-dithiane ring. The 2-position of the ring is connected via a (CH₂)₃ chain to the nitrogen of a sulfonamide group. This sulfonamide group is linked to a 3,5-dimethylphenyl ring. The 5-position of the ring is connected via another (CH₂)₃ chain to the nitrogen of a second sulfonamide group, which is also linked to a 3,5-dimethylphenyl ring. The 6-position of the ring is connected via a (CH₂)₃ chain to the nitrogen of a third sulfonamide group, which is linked to a 3,5-dimethylphenyl ring. The 3-position of the ring is connected via a (CH₂)₃ chain to the nitrogen of a fourth sulfonamide group, which is linked to a 3,5-dimethylphenyl ring. The sulfonamide groups are represented as -SO₂-N(Et)-, where Et is an ethyl group. The 3,5-dimethylphenyl groups are represented as a benzene ring with methyl groups at the 3 and 5 positions.

$$\begin{array}{c} | \\ \text{Me} \end{array}$$

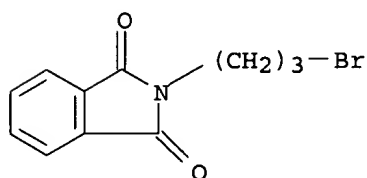
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

Relative stereochemistry.



●2 Cl⁻

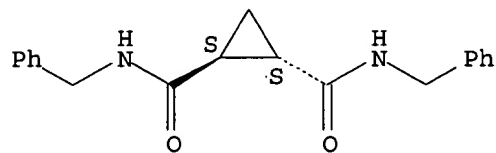
L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1H-Isoindole-1,3(2H)-dione, 2-(3-bromopropyl)- (9CI)
 MF C11 H10 Br N O2
 CI COM



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedicarboxamide, N,N'-bis(phenylmethyl)-, (1R,2R)-rel- (9CI)
 MF C19 H20 N2 O2

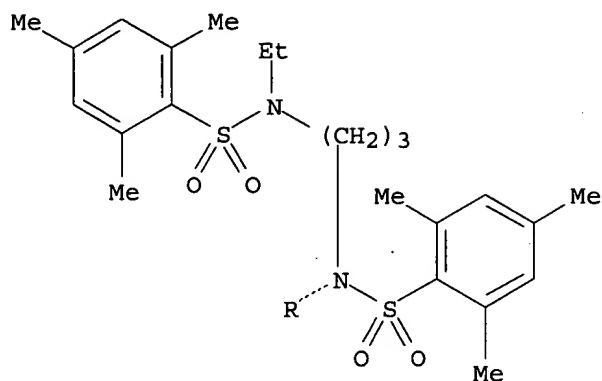
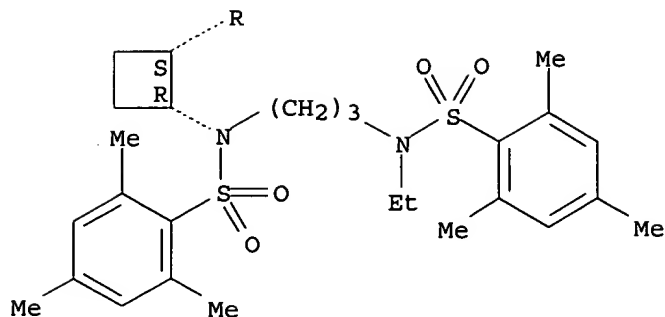
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonamide, N,N'-(1R,2S)-1,2-cyclobutanediylbis[N-[3-[ethyl[(2,4,6-trimethylphenyl)sulfonyl]amino]propyl]-2,4,6-trimethyl-, rel- (9CI)
 MF C50 H72 N4 O8 S4

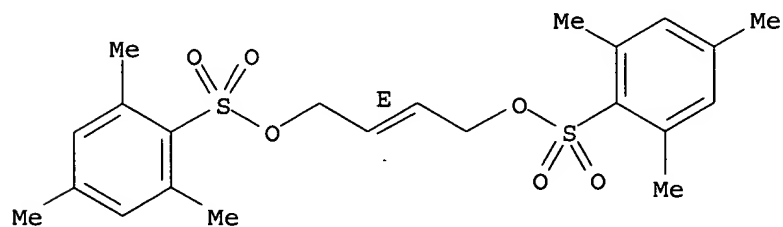
Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN Benzenesulfonic acid, 2,4,6-trimethyl-, (2E)-2-butene-1,4-diyl ester (9CI)
 MF C22 H28 O6 S2

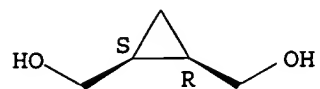
Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 69 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN 1,2-Cyclopropanedimethanol, (1R,2S)-rel- (9CI)
 MF C5 H10 O2

Relative stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED



US006392098B1

(12) **United States Patent**
Frydman et al.(10) **Patent No.:** US 6,392,098 B1
(45) **Date of Patent:** May 21, 2002(54) **CONFORMATIONALLY RESTRICTED
POLYAMINES**(75) **Inventors:** Benjamin J. Frydman, Madison;
Laurence J. Marton, Fitchburg;
Venodhar K. Reddy, Madison;
Aldonla L. Valasinas, Madison;
Donald T. Witiak, Madison, all of WI
(US)(73) **Assignee:** Wisconsin Alumni Research
Foundation, Madison, WI (US)(*) **Notice:** Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 0 days.(21) **Appl. No.:** 09/280,278(22) **Filed:** Mar. 29, 1999**Related U.S. Application Data**(63) Continuation of application No. 08/951,015, filed on Oct.
15, 1997, now Pat. No. 5,889,061.(51) **Int. Cl.⁷** A61K 31/13(52) **U.S. Cl.** 564/509; 564/510; 514/674(58) **Field of Search** 514/674; 564/509,
564/510(56) **References Cited****U.S. PATENT DOCUMENTS**

5,889,061 A * 3/1999 Frydman et al.

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Spermine and Its Cytotoxic Analogs on Nucleosome Forma-
tion on Relaxed and Supercoiled DNA In Vitro, *Biochem-*
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Primary Examiner—Samuel Barts(74) *Attorney, Agent, or Firm*—Joseph T. Leone, Esq.;
DeWitt Ross & Stevens S.C.(57) **ABSTRACT**

Compounds of Formula I:

wherein A is C₂-C₆ alkene, C₃-C₆ cycloalkyl, cycloalkenyl,
or cycloaryl; B is independently a single bond, C₁-C₆ alkyl
alkenyl; D is independently C₁-C₆ alkyl or alkenyl, or
C₃-C₆ cycloalkyl, cycloalkenyl, or cycloaryl; and E is
independently H, C₃-C₆ alkyl or alkenyl; and
pharmaceutically-suitable salts thereof; a synthetic method
therefor, pharmaceutical dosage forms containing one of
more of these compounds, and use of these compounds in
the treatment of neoplastic cell growth, are disclosed.

22 Claims, 20 Drawing Sheets

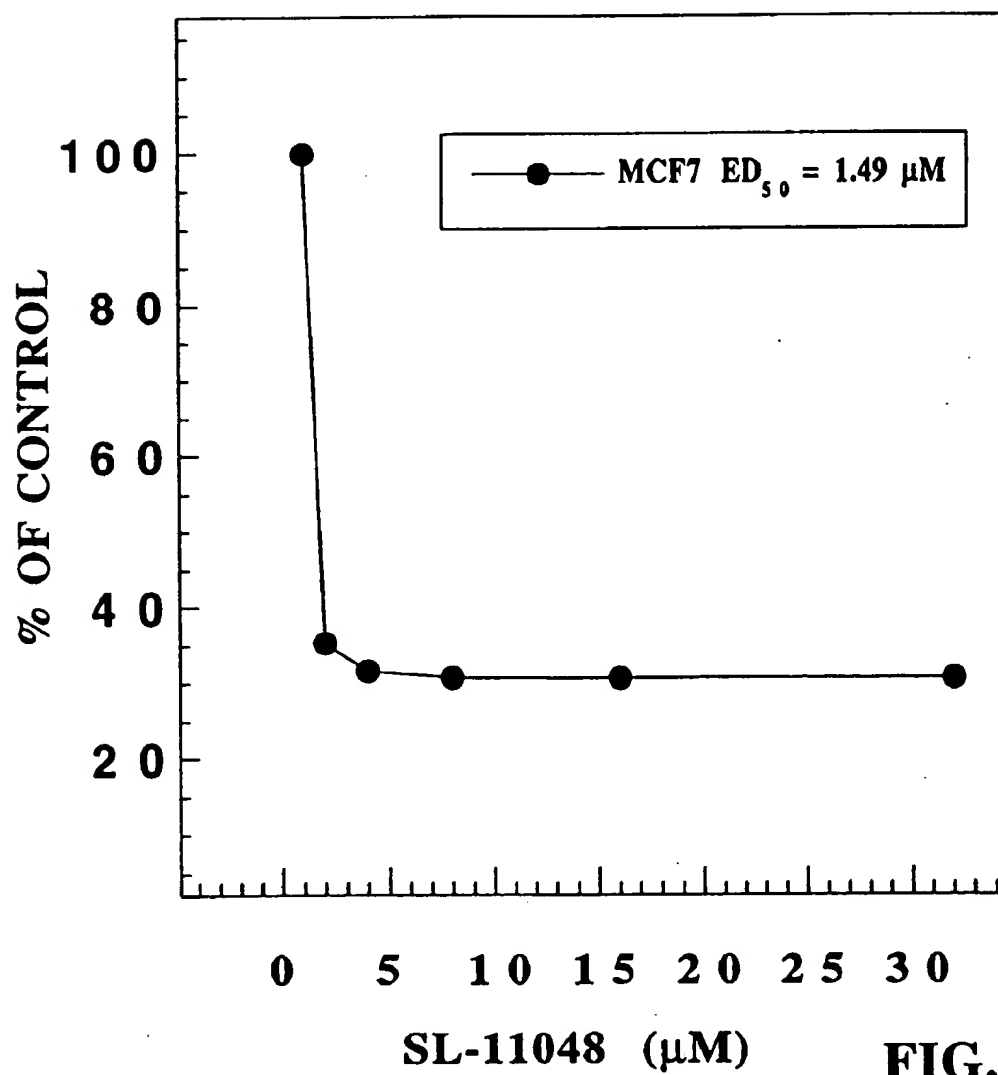


FIG. 1

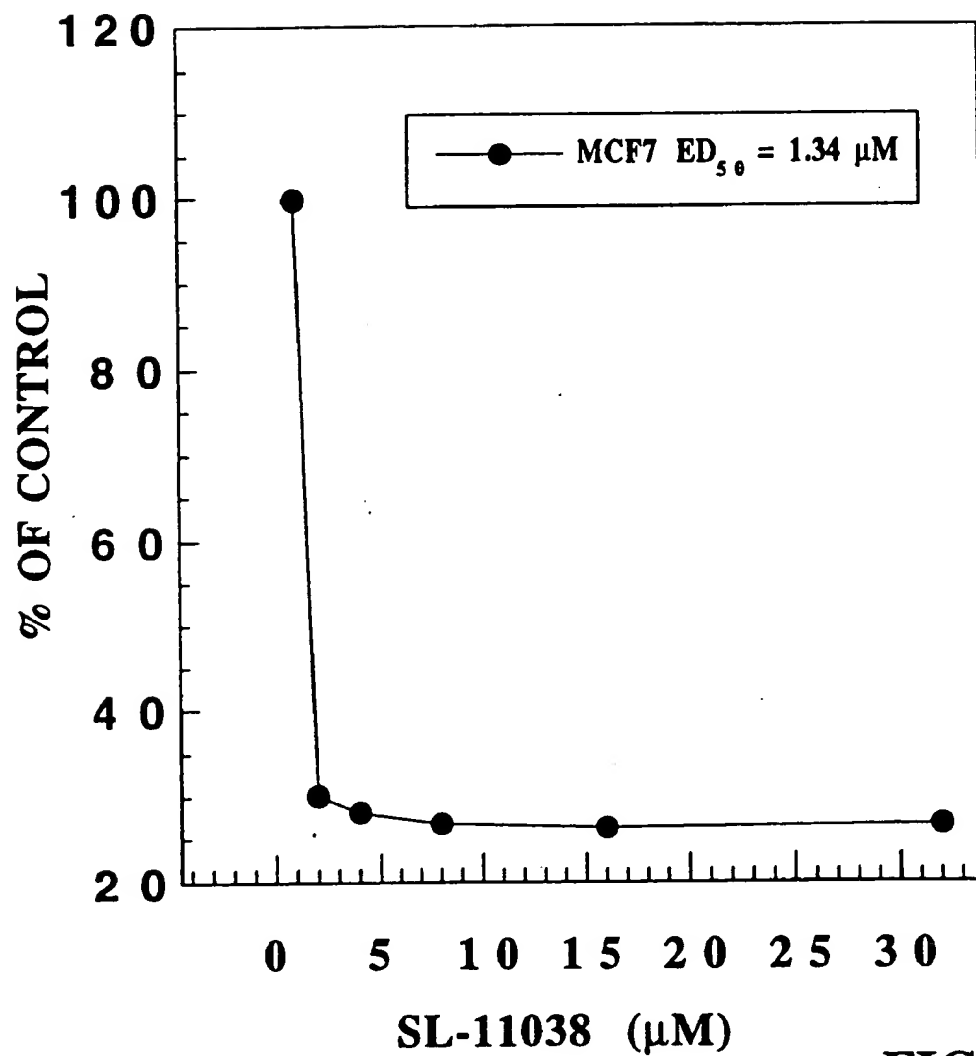
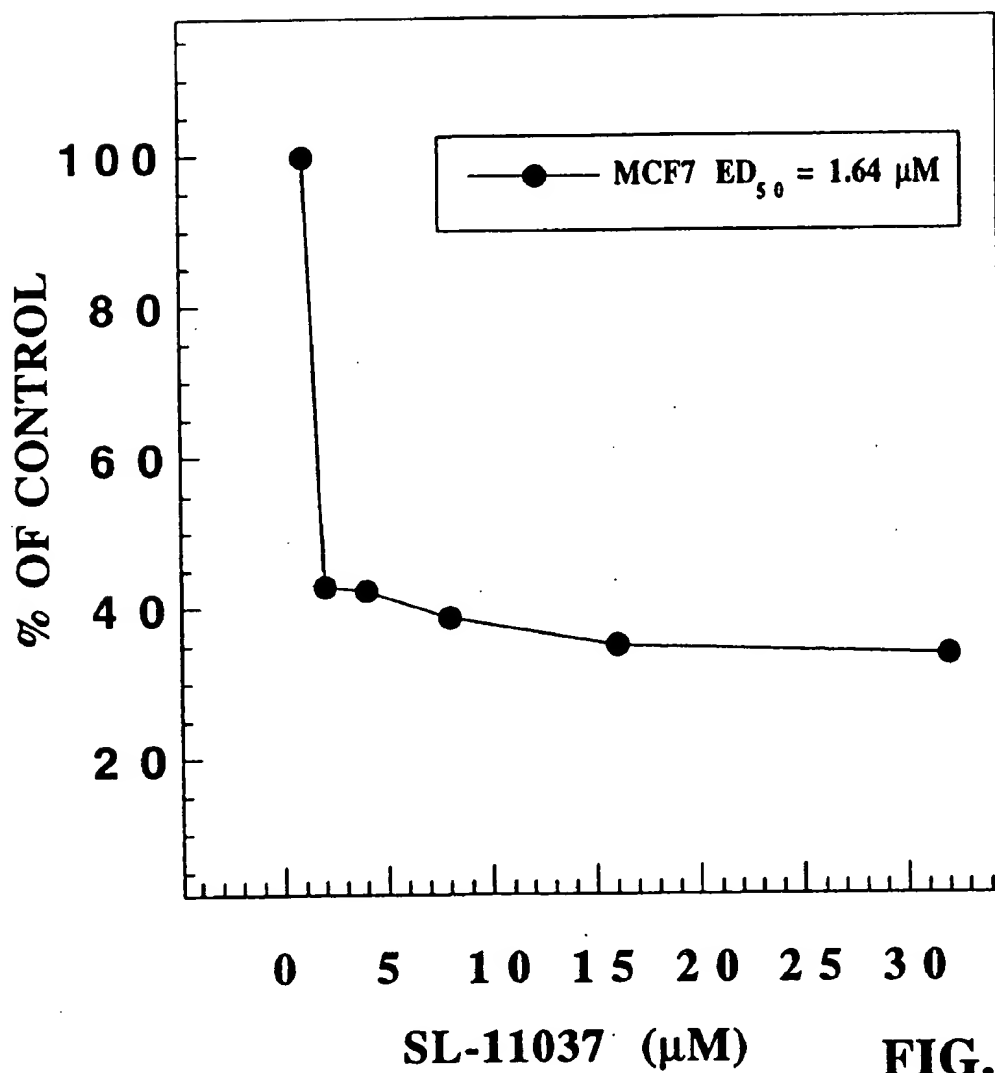


FIG. 2



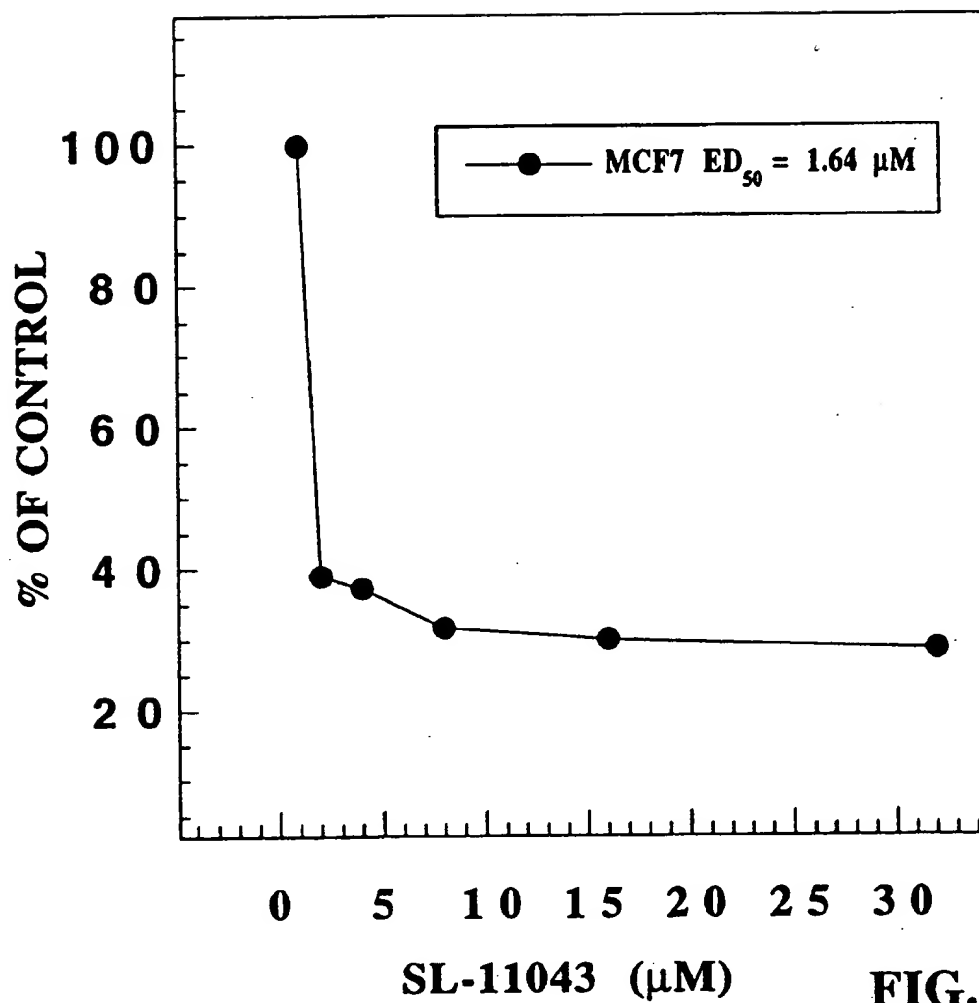
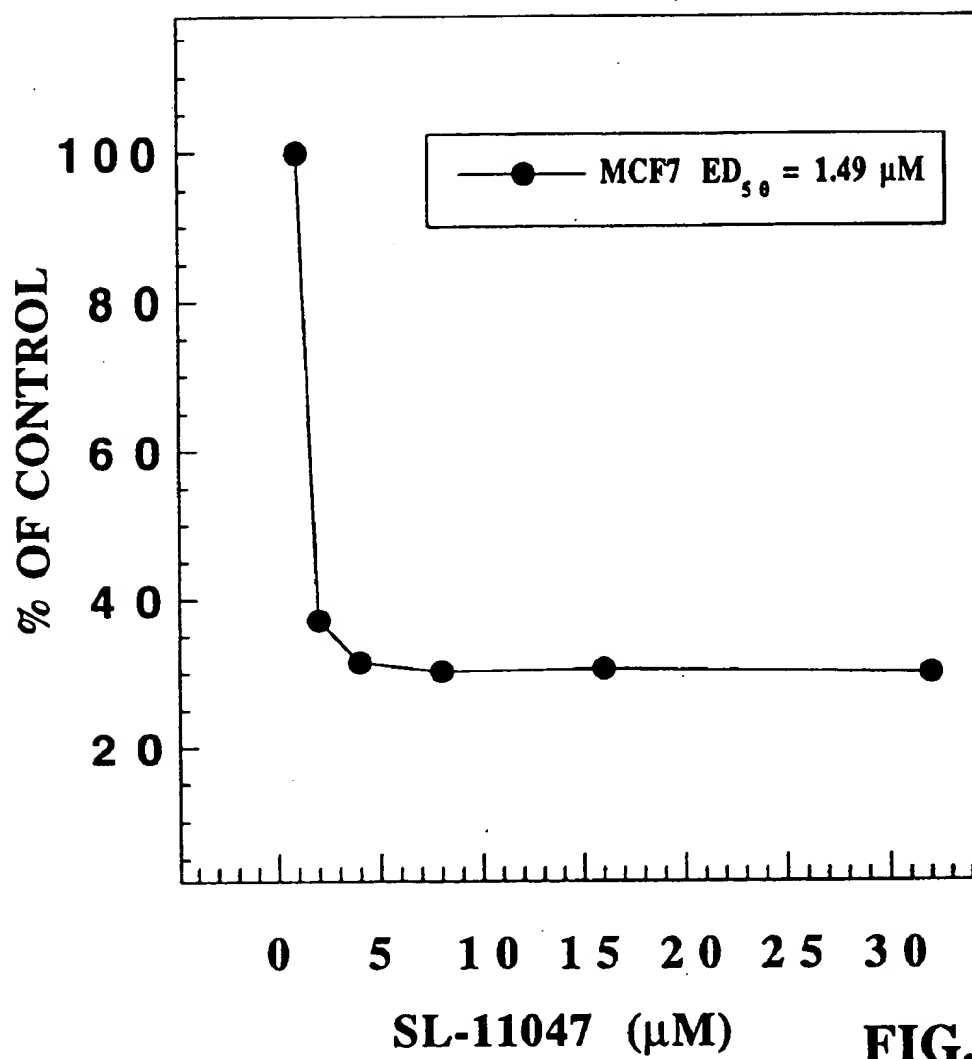


FIG. 4



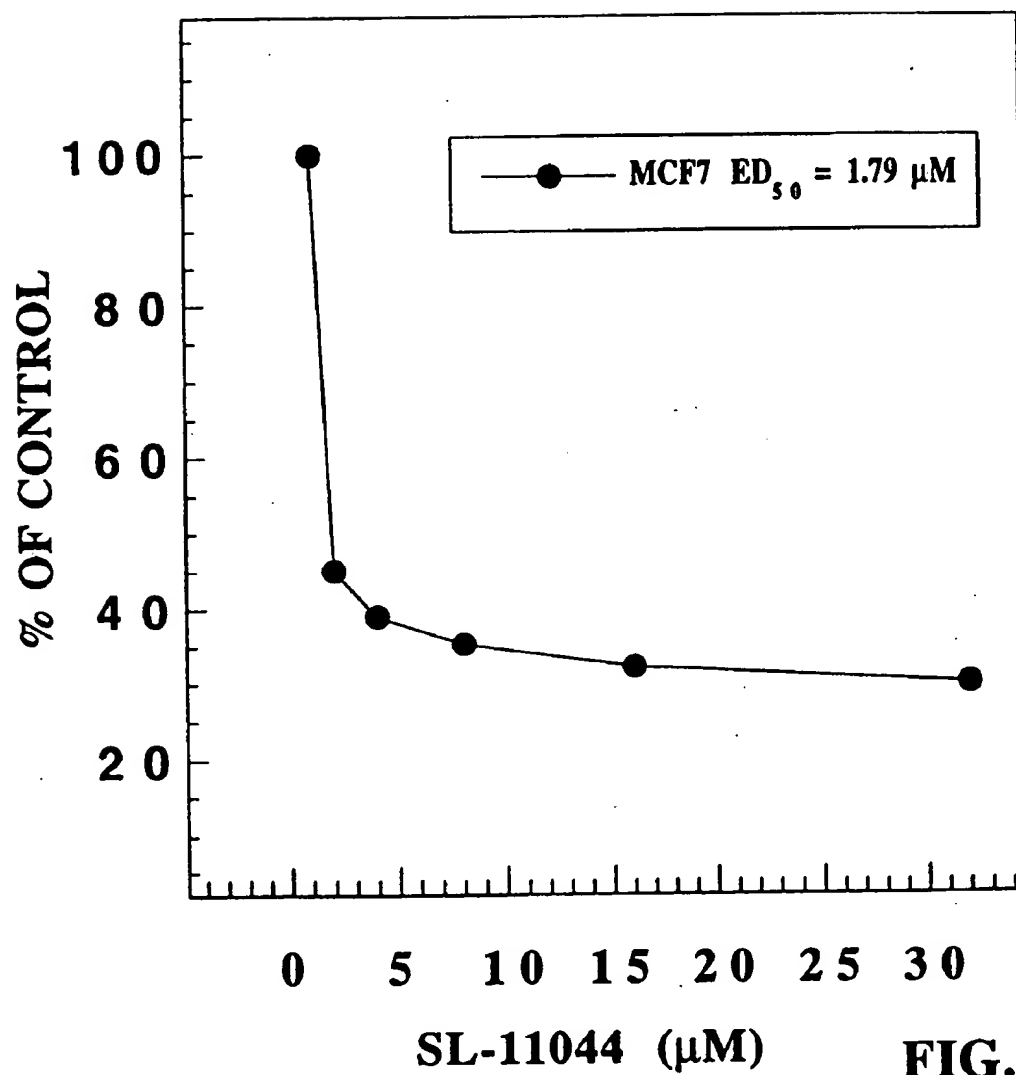


FIG. 6

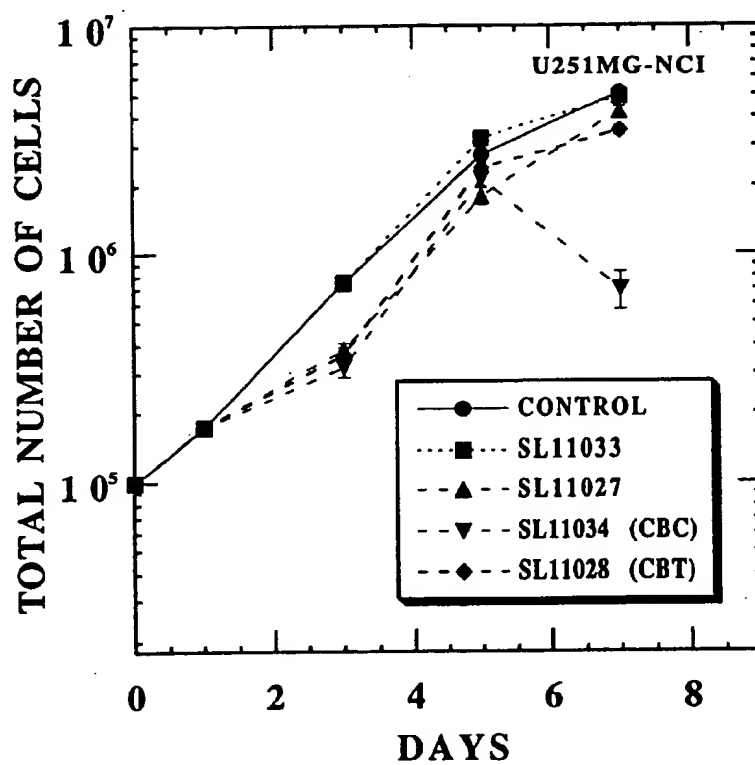
EFFECT OF 4 DIFF. DRUGS (10 μ M) ON U251MG-NCI CELLS

FIG. 7

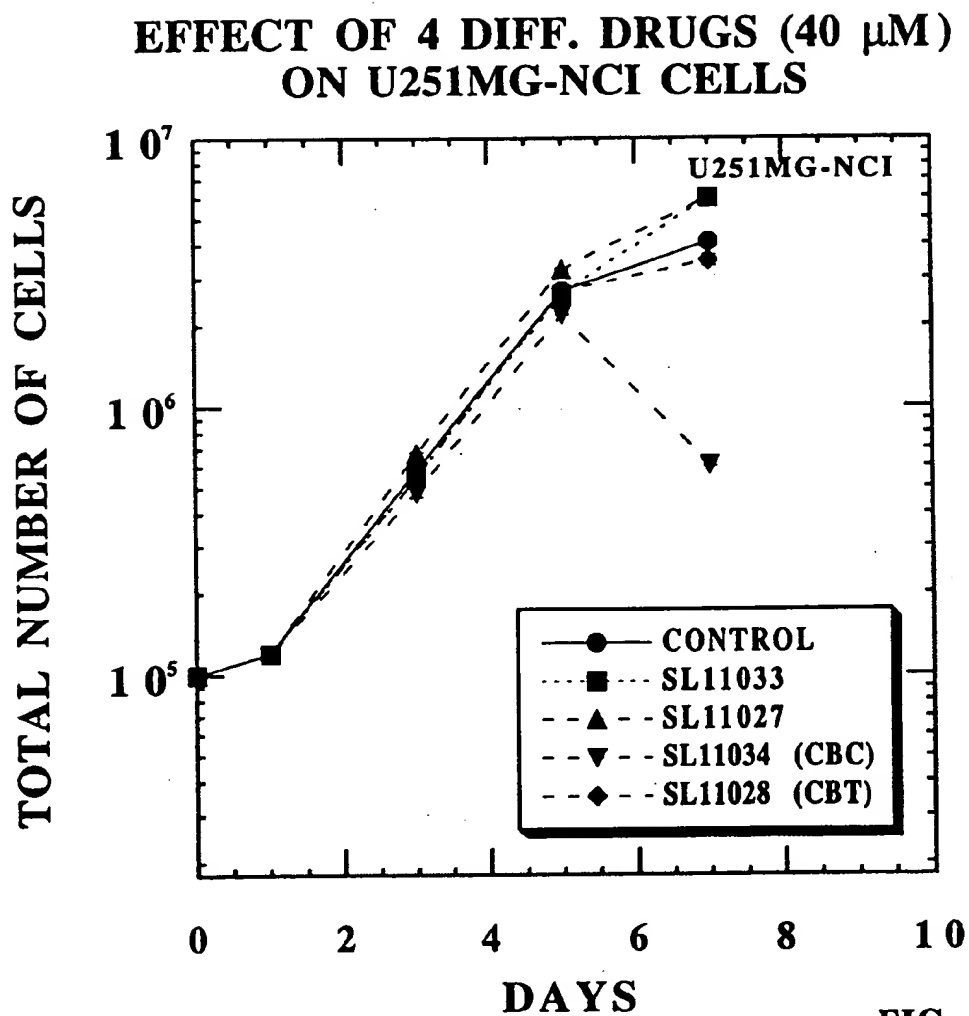


FIG. 8

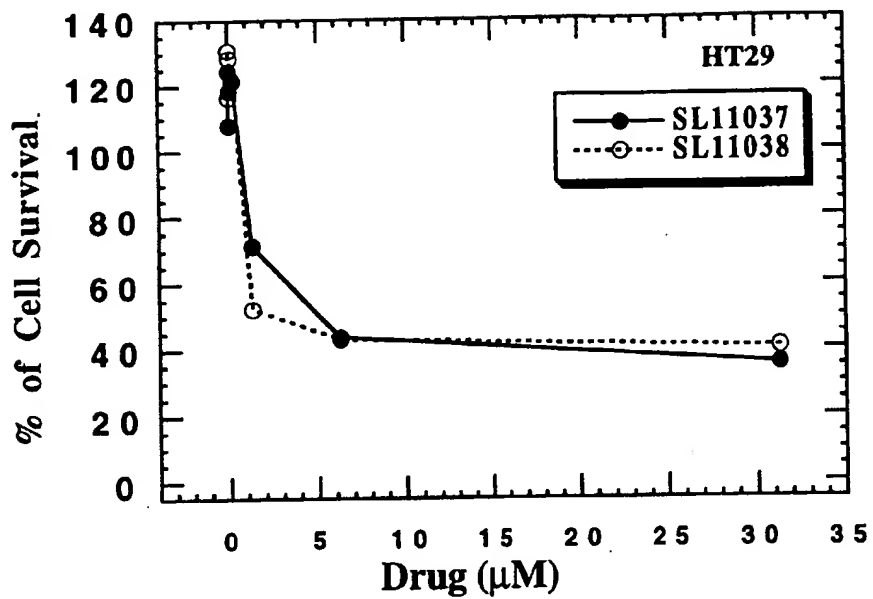


Fig. 9A

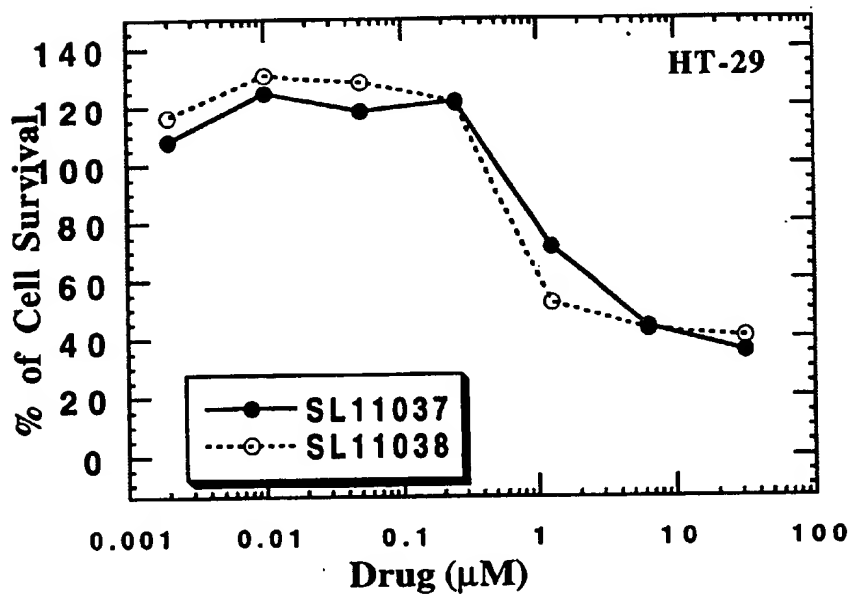


Fig. 9B

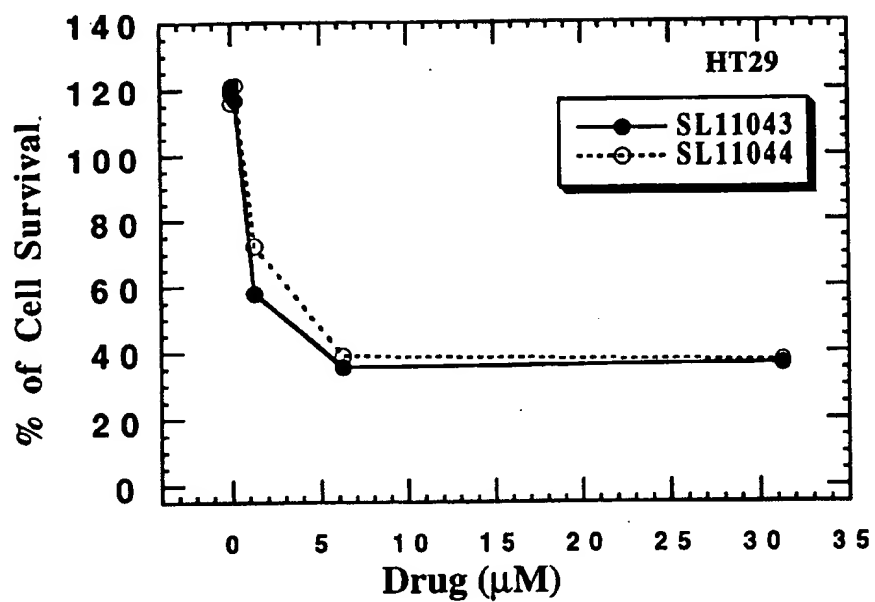


Fig. 10A

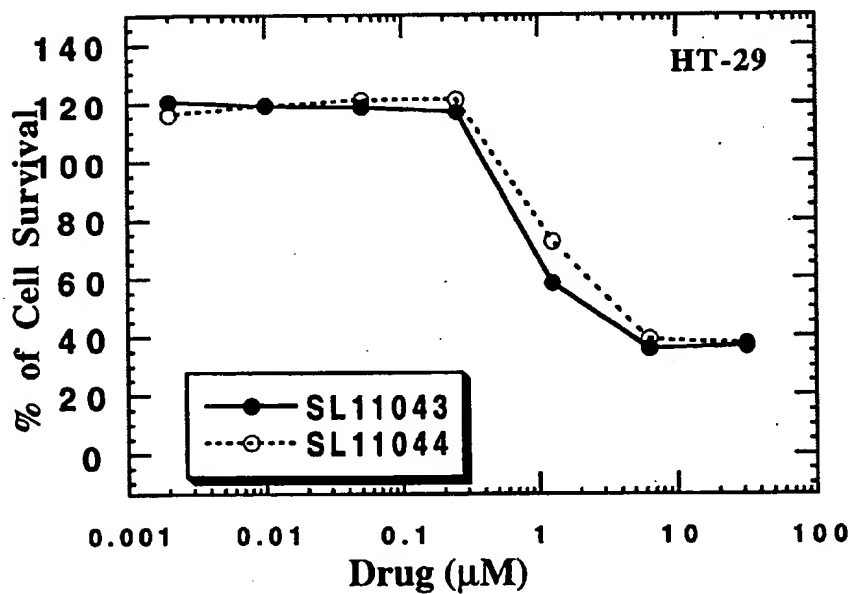


Fig. 10B

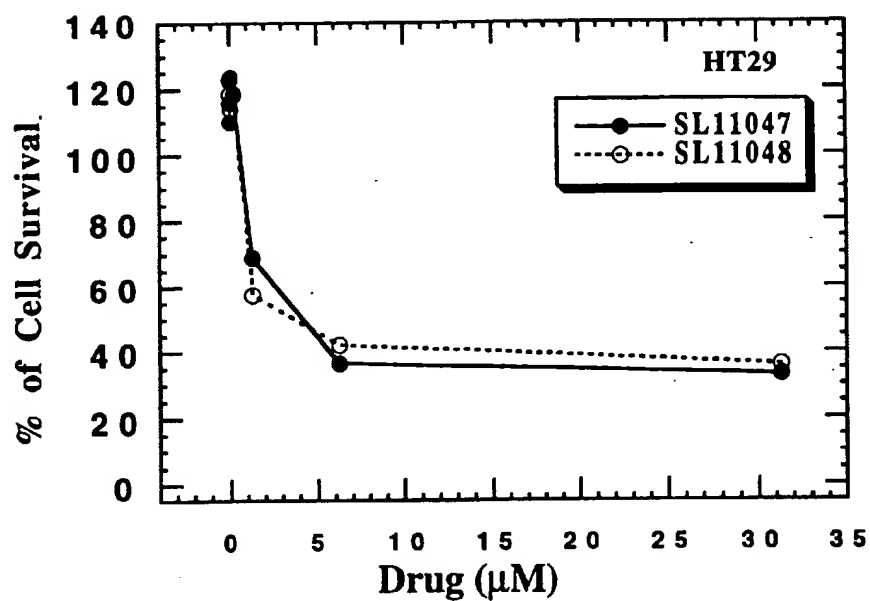


Fig. 11A

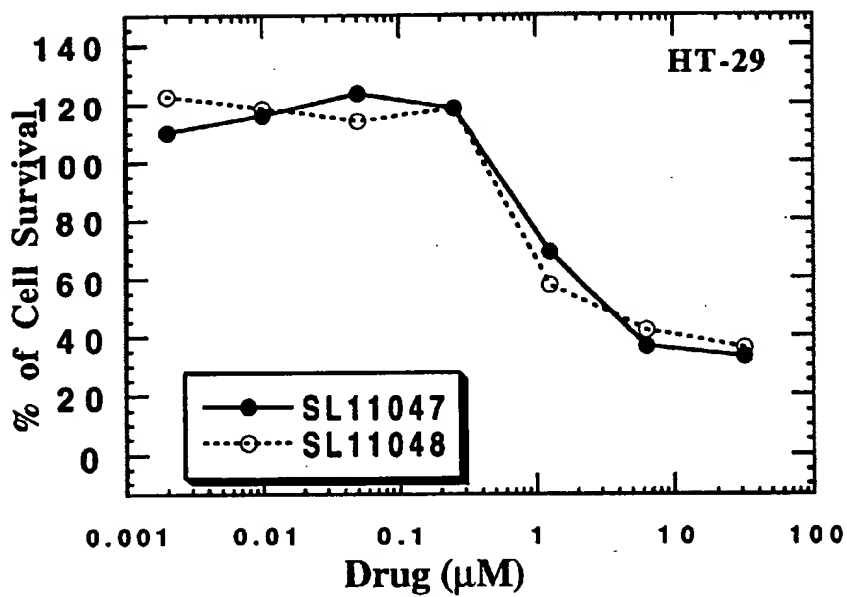


Fig. 11B

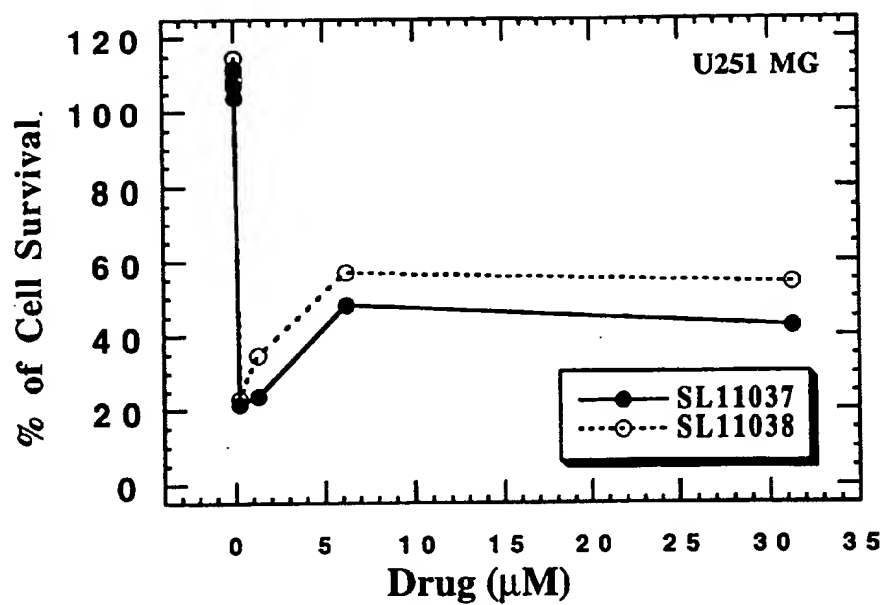


Fig. 12A

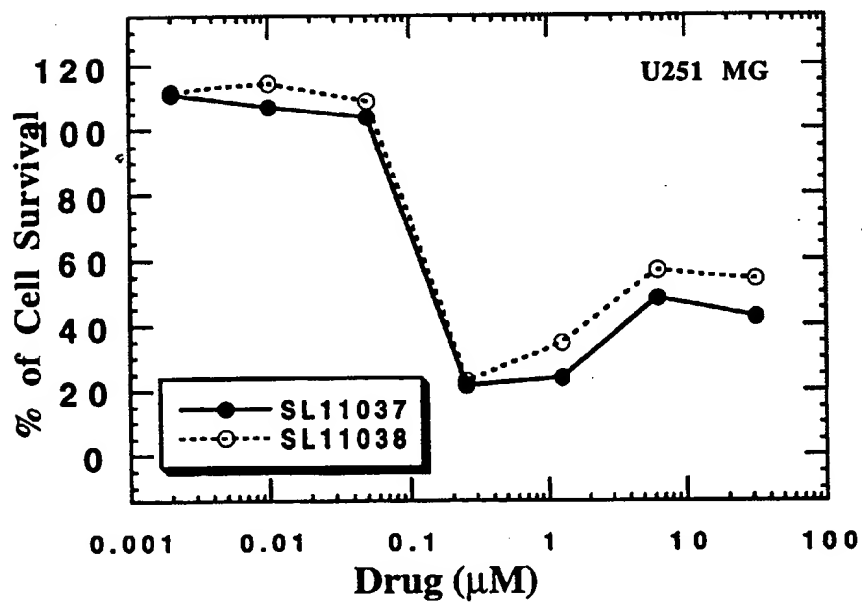


Fig. 12B

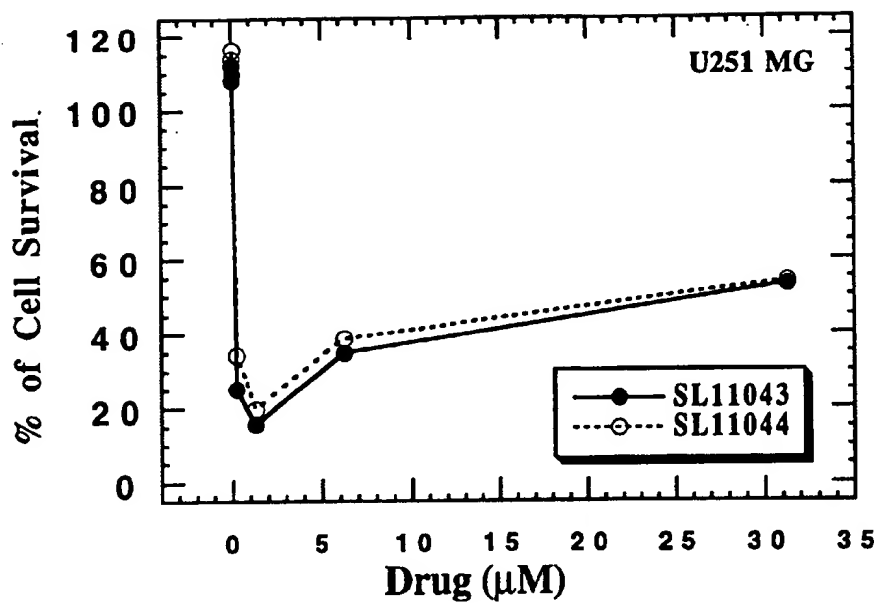


Fig. 13A

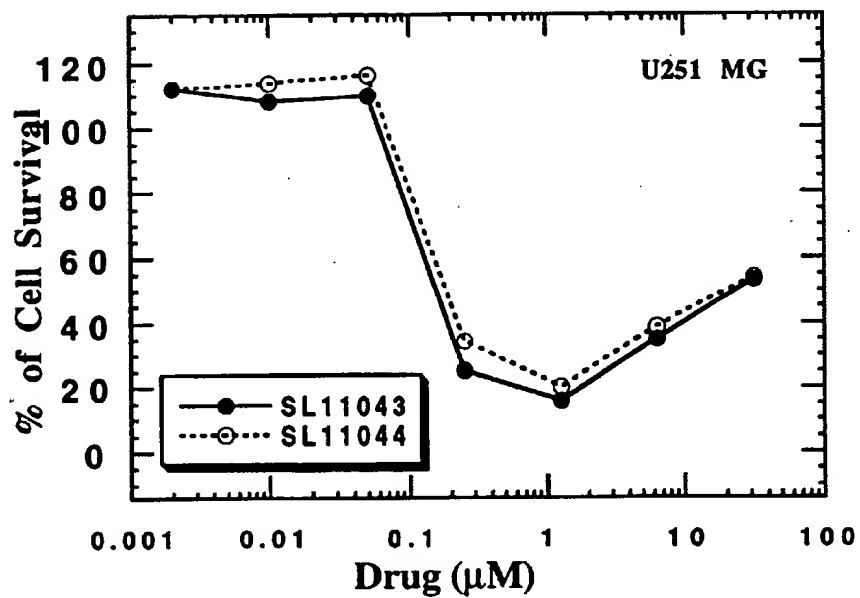


Fig. 13B

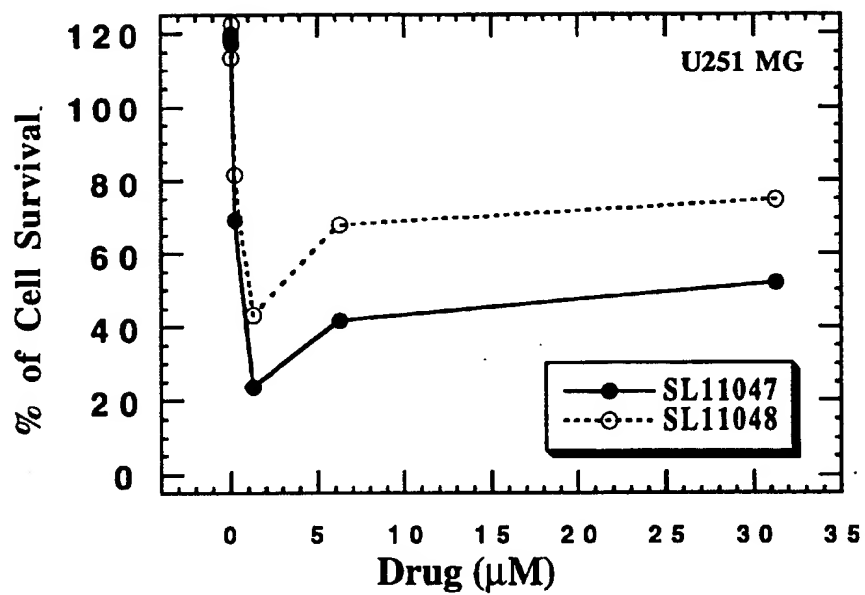


Fig. 14A

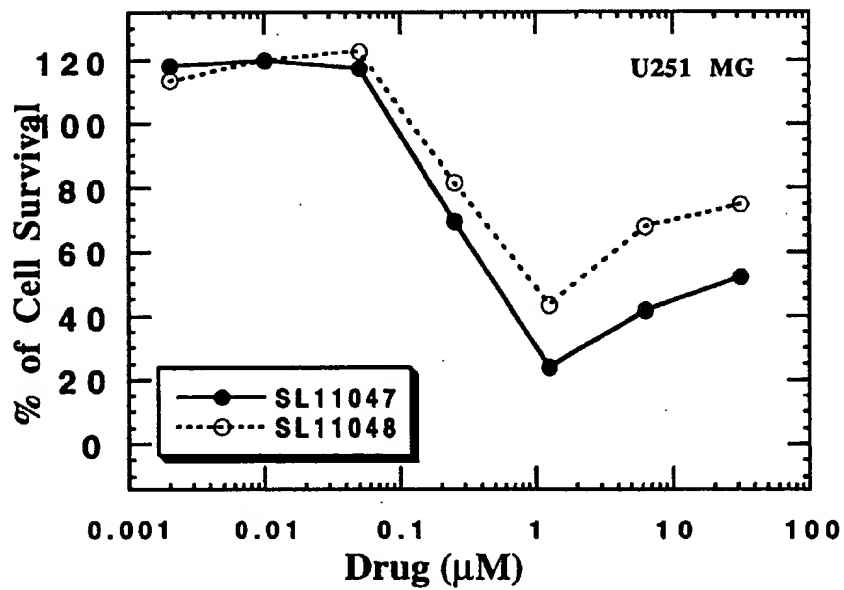


Fig. 14B

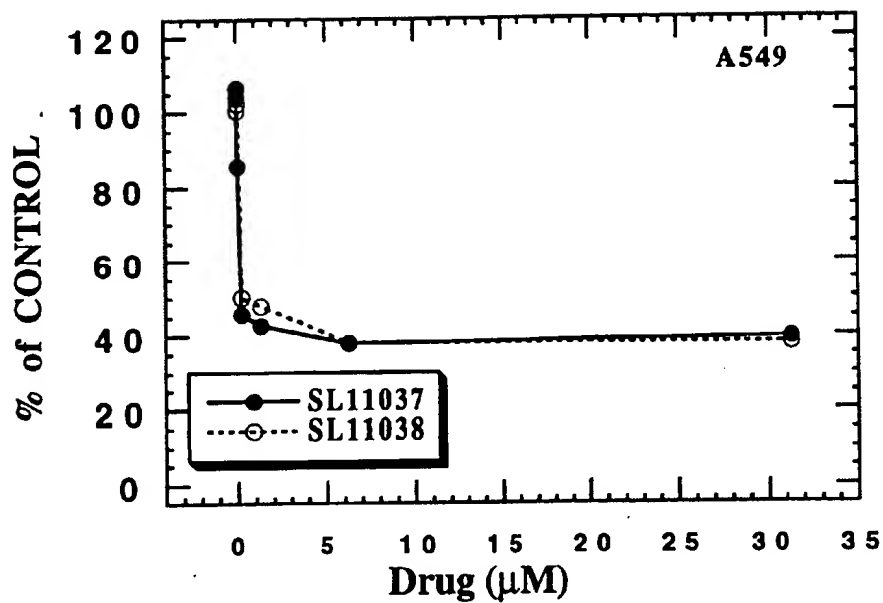


Fig. 15A

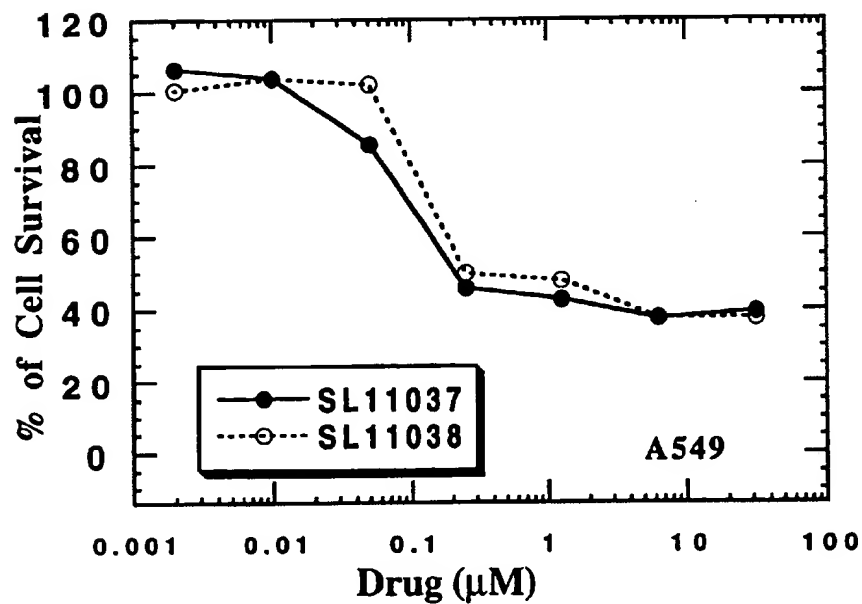


Fig. 15B

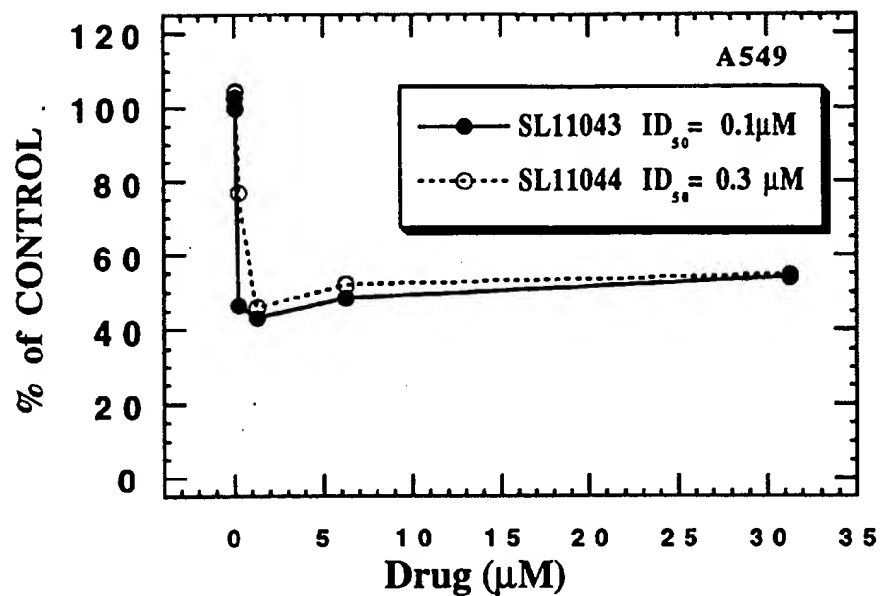


Fig. 16A

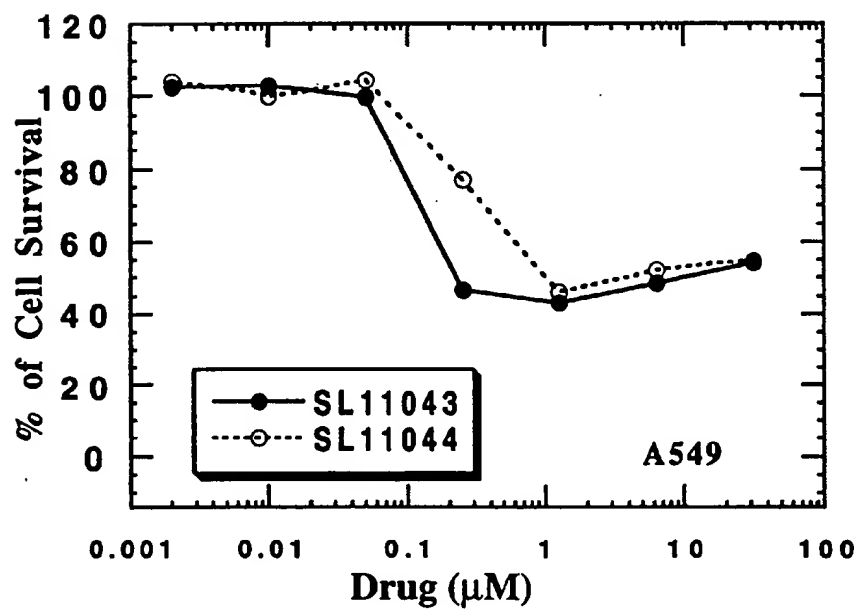


Fig. 16B

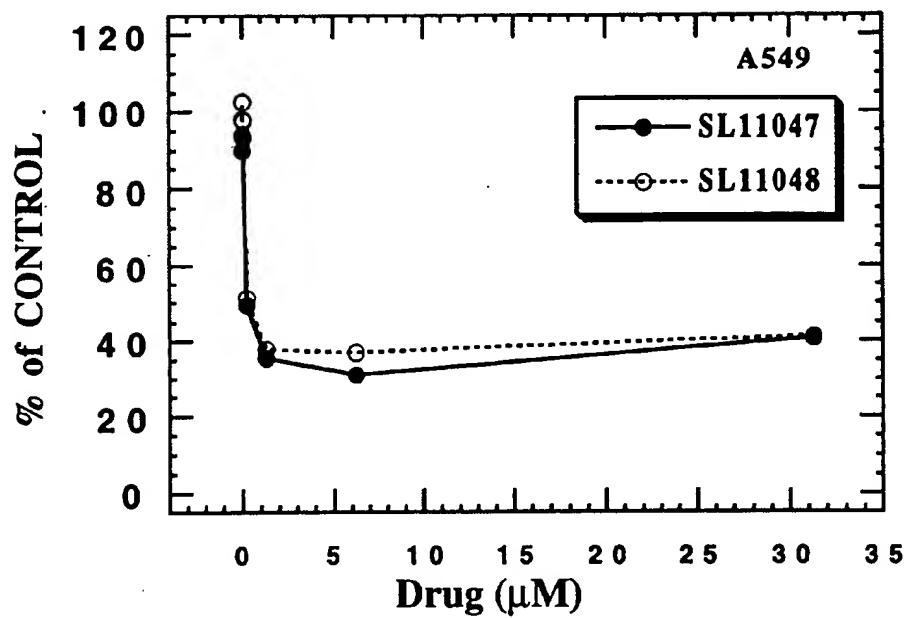


Fig. 17A

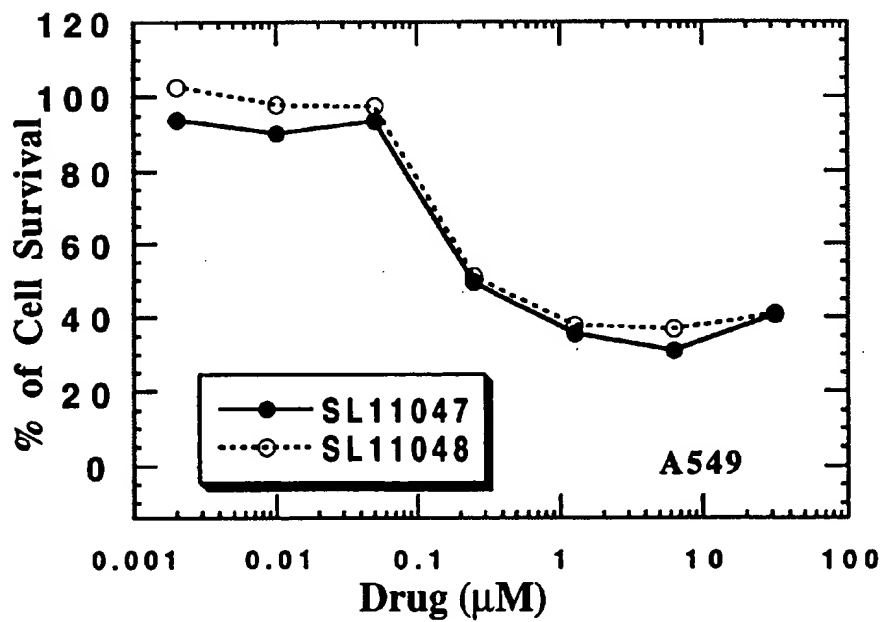


Fig. 17B

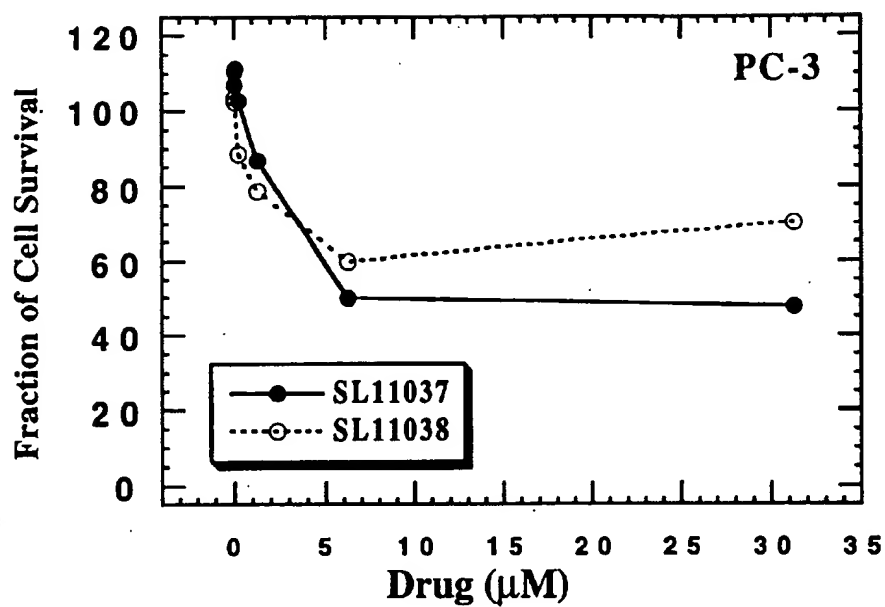


Fig. 18A

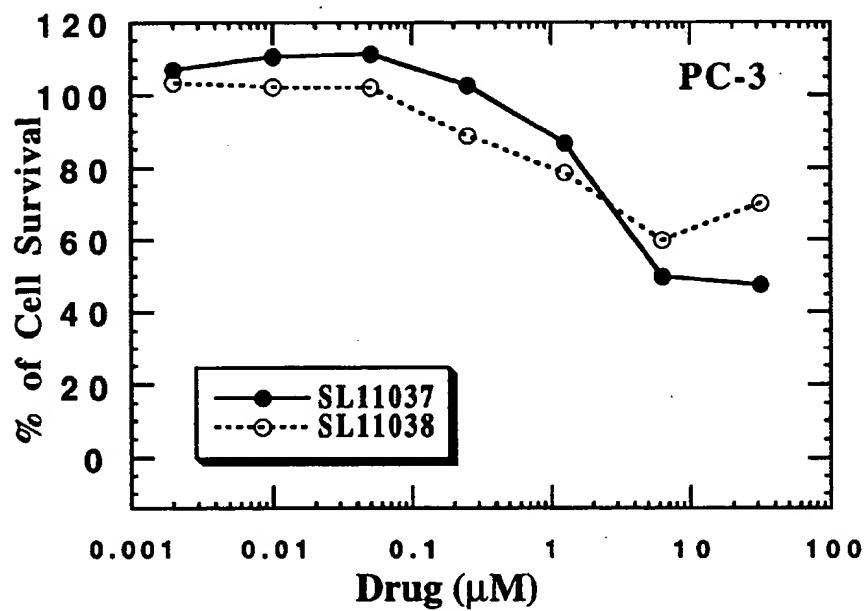


Fig. 18B

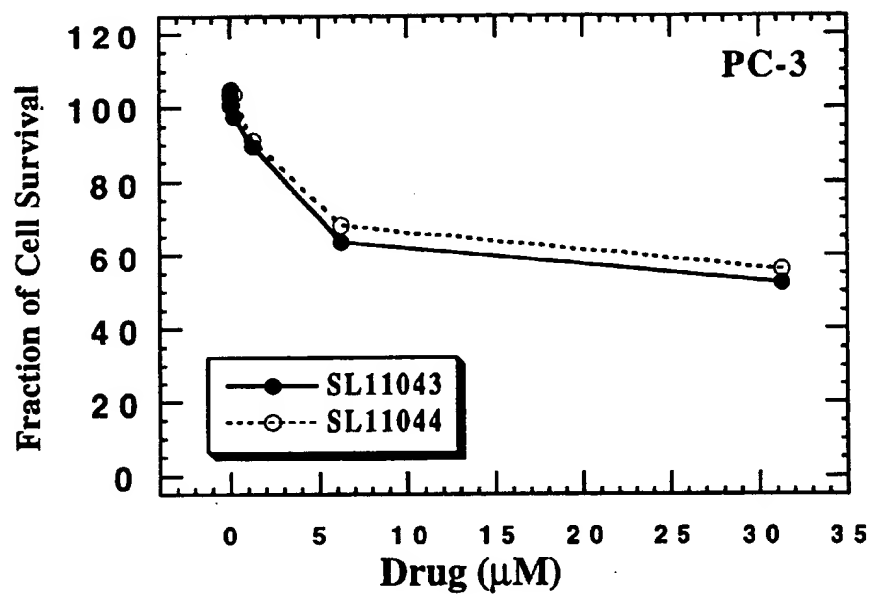


Fig. 19A

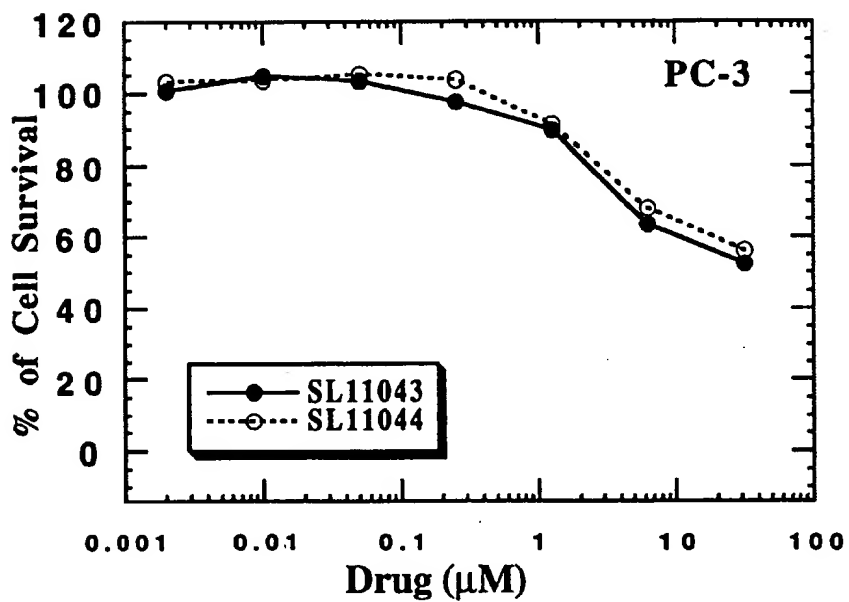


Fig. 19B

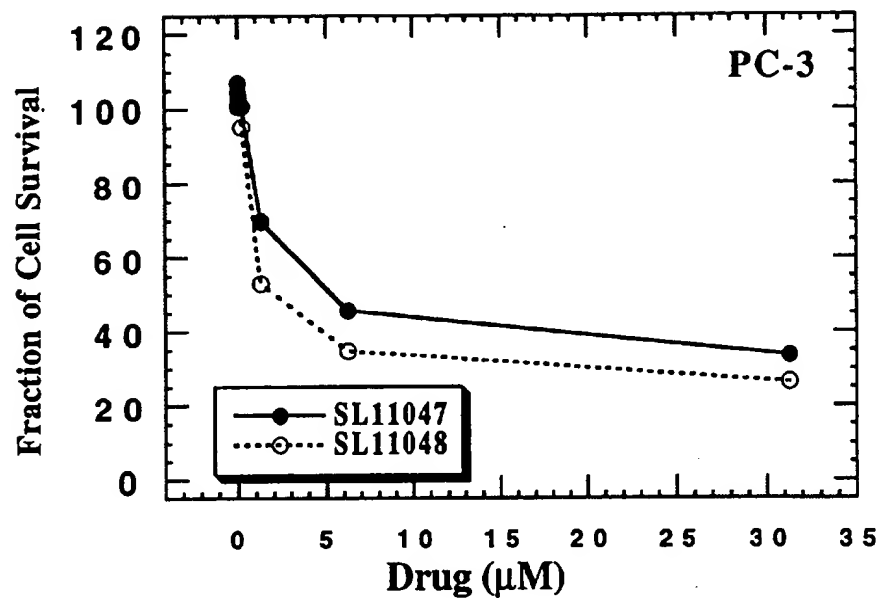


Fig. 20A

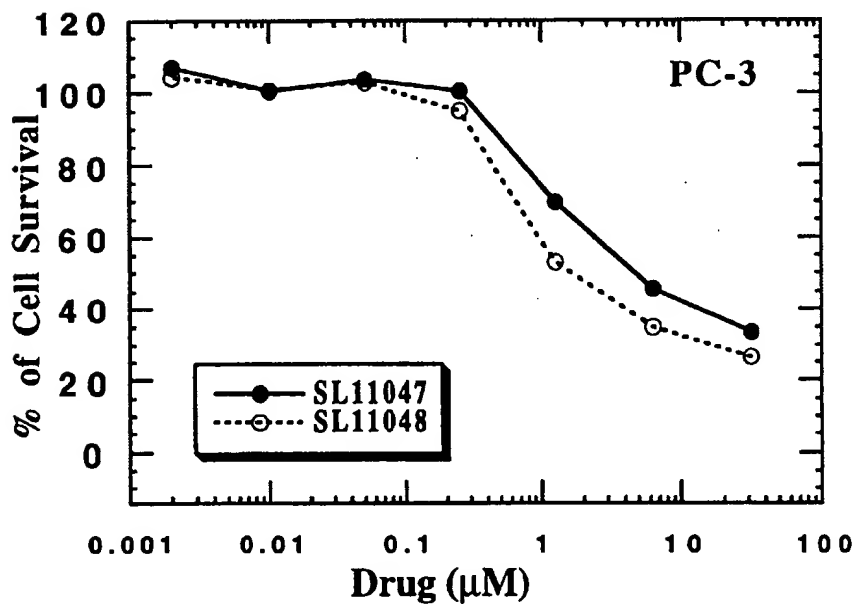


Fig. 20B

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CONFORMATIONALLY RESTRICTED POLYAMINES

This is a continuation of application Ser. No. 08/951,015, filed on Oct. 15, 1997 now U.S. Pat. No. 5,889,061.

FIELD OF THE INVENTION

The present invention is directed to novel conformationally restricted polyamines and their use in the selective inhibition of neoplastic cell growth.

BIBLIOGRAPHY

Complete bibliographic citations to the references mentioned below are listed in the Bibliography, immediately preceding the claims. All of the references cited below are incorporated herein by reference in their entirety.

DESCRIPTION OF THE PRIOR ART

It has been known since the 1950's that conformation is a determinant in the spatial arrangement of functional groups, and that enzymes or drug receptors prefer specific ligand conformations or a specific distribution of conformations. This fruitful concept led to many decisive successes in drug design. A few examples will suffice to illuminate the subject.

The synthesis of conformationally constrained analogs of an inherently conformationally flexible substance such as acetyl choline helped to secure its "bioactive conformations," i.e., those conformers which are active at the muscarinic and nicotinic receptors. The trans-cyclopropyl analog of acetyl choline was shown to be preferred by the muscarinic receptor. Conformationally restricted analogs of dopamine, GABA, glutamic acid, histamine and serotonin have been obtained by introducing rigid rings into their structures. The constrained analogs have valuable chemotherapeutic effects.

The use of conformational restriction has also been very fruitful in the design of bioactive polypeptides. Polypeptides have so many flexible torsion angles that enormous numbers of conformations are possible in solution. The introduction of rings into the linear peptide chains reduces the number of conformations and has allowed the preparation of several biologically active substances. For instance, a cyclic hexapeptide possessing somatostatin activity is known.

Conformationally restricted enkephalin analogs are known, as are bicyclic lactam inhibitors (enalapril and enalaprilat) of the angiotensin converting enzyme.

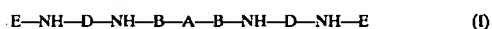
Similar strategies have recently led to the development of a peptidomimetic benzodiazepine containing at least two conformational restrictions: a bicyclic heterocycle and an acetylene linker. The benzodiazepine is a non-peptide RGD (Arg-Gly-Asp) receptor antagonist.

The concept of conformational restriction led to the discovery that the bioactive conformation of the immunosuppressor cyclosporin A (CsA) only binds to cyclophilin A when the amide bond between the 9-position and 10-position residues in CsA is trans.

However, the prior art is silent regarding biologically active polyamines which are conformationally restricted by the introduction of one or more ring structures into the polyamine skeleton.

SUMMARY OF THE INVENTION

The present invention is directed to conformationally restricted polyamines of Formula I:



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wherein A is selected from the group consisting of C₂-C₆ alkene and C₃-C₆ cycloalkyl, cycloalkenyl, and cycloaryl; B is independently selected from the group consisting of a single bond and C₁-C₆ alkyl and alkenyl; D is independently selected from the group consisting of C₁-C₆ alkyl and alkenyl, and C₃-C₆ cycloalkyl, cycloalkenyl, and cycloaryl; E is independently selected from the group consisting of H, C₁-C₆ alkyl and alkenyl; and pharmaceutically-suitable salts thereof.

The invention is also drawn to a method of synthesizing the Formula I compounds. Here, a compound of Formula II:

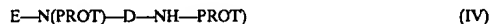


is reacted with with a protecting reagent, preferably mesitylenesulfonyl chloride, to yield a compound of Formula III:

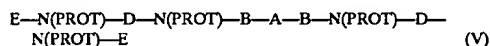


wherein PROT is the protecting group.

Then, the Formula III compound is reacted with a compound of Formula IV:



to yield a compound of Formula V:



It is much preferred that the protecting group, PROT, in both the Formula III intermediate and the Formula IV intermediate be a mesitylenesulfonyl moiety.

The Formula V compound is then deprotected to yield a compound of Formula I.

The present invention is also drawn to pharmaceutical unit dosage forms containing one or more compounds of Formula I as described above in combination with a pharmaceutically-suitable carrier.

The present invention is also drawn to use of these novel conformationally restricted polyamines. These polyamines have utility as potent antineoplastic agents for use in mammals, including humans.

They also have utility as shape-restricted probes for the study of active-site geometry for enzyme and/or DNA interactions. Because the backbone of these molecules is conformationally restricted, they can only assume a very limited number of 3-dimensional shapes. By assessing the binding of these compounds to various enzyme active sites, or the their ability to interact with DNA, insight is gained into the particular spacial geometry required of enzyme agonists, antagonists and DNA binders. Both the compounds per se and their use is novel.

The present invention was inspired by a study of past failures in the use of polyamine compounds as chemotherapeutic agents in the treatment antiproliferative diseases. It is known that the polyamines spermidine and spermine are essential for normal cell growth. Potent antiproliferative agents have been developed which interfere with the biosynthesis of these compounds, thereby preventing cell proliferation. One of these agents, difluoromethylornithine (DFMO), is currently being studied in humans as a chemopreventive.

The therapeutic success of DFMO is, however, significantly marred by cellular uptake of exogenous polyamines via the polyamine transport system. Cellular uptake of polyamines from the extracellular milieu compensates for the endogenous depletion of the cellular polyamine pools due to the effects of DFMO. Since most food is rich in

polyamines (e.g., 100 mL of orange juice contains approximately 400 ppm of putrescine, the spermine precursor), the antineoplastic effect of synthetic analogues which act by depleting the endogenous pool of polyamines is greatly diminished.

The present invention introduces a new approach for the use of polyamine analogs as agents for the treatment of cancer. Notably, both spermidine and spermine interact with DNA. These interactions induce structural changes in isolated DNA. Computer modeling and physico-chemical studies indicate that spermine induces conformational changes in defined DNA sequences. The presently described analogs of natural polyamines may interact with DNA differently than does spermine, resulting in the inhibition of tumor cell growth in culture.

In short, while not being bound to a particular mode of action, the present invention attacks the problem of unchecked cell proliferation by presumably changing DNA conformation via a polyamine analog/DNA interaction. The novel, conformationally restricted polyamines of the present invention display potent anti-proliferative activity.

Consequently, it is the principal aim of the present invention to provide novel, conformationally restricted compounds for use in the treatment of neoplastic cell growth.

These and other aims, objects, and advantages of the present invention will become clear upon a complete reading of the Detailed Description and claims, below.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a graph depicting the in vitro effect of increasing concentrations of SL-11048 (Compound 57) on the survival of cultured breast cancer cells MCF7. $ED_{50}=1.49 \mu M$.

FIG. 2 is a graph depicting the in vitro effect of increasing concentrations of SL-11038 (Compound 23) on the survival of cultured breast cancer cells MCF7. $ED_{50}=1.34 \mu M$.

FIG. 3 is a graph depicting the in vitro effect of increasing concentrations of SL-11037 (Compound 28) on the survival of cultured breast cancer cells MCF7. $ED_{50}=1.64 \mu M$.

FIG. 4 is a graph depicting the in vitro effect of increasing concentrations of SL-11043 (Compound 48) on the survival of cultured breast cancer cells MCF7. $ED_{50}=1.64 \mu M$.

FIG. 5 is a graph depicting the in vitro effect of increasing concentrations of SL-11047 (Compound 58) on the survival of cultured breast cancer cells MCP7. $ED_{50}=1.49 \mu M$.

FIG. 6 is a graph depicting the in vitro effect of increasing concentrations of SL-11044 (Compound 47) on the survival of cultured breast cancer cells MCF7. $ED_{50}=1.79 \mu M$.

FIG. 7 is a graph depicting the in vitro effect of a 10 μM concentration of SL-11033 (13, ■), SL-11027 (12, ▲), SL-11034 (36, ▼), and SL-11028 (35, ◆) on the growth of brain cancer cells U251MG-NCI cells. Control=●.

FIG. 8 is a graph depicting the in vitro effect of a 40 μM concentration of SL-11033 (13, ■), SL-11027 (12, ▲), SL-11034 (36, 569), and SL-11028 (35, ◆) on the growth of brain cancer cells U251MG-NCI cells. Control=●.

FIGS. 9A and 9B are graphs depicting the in vitro effect of increasing concentrations of SL-11037 (Compound 28, ●) and SL-11038 (Compound 23, ○) on the survival of cultured human colon cancer cells HT-29.

FIGS. 10A and 10B are graphs depicting the in vitro effect of increasing concentrations of SL-11043 (Compound 48,

●) and SL-11044 (Compound 47, ○) on the survival of cultured human colon cancer cells HT-29.

FIGS. 11A and 11B are graphs depicting the in vitro effect of increasing concentrations of SL-11047 (Compound 58, ●) and SL-11048 (Compound 57, ○) on the survival of cultured human colon cancer cells HT-29.

FIGS. 12A and 12B are graphs depicting the in vitro effect of increasing concentrations of SL-11037 (Compound 28, ●) and SL-11038 (Compound 23, ○) on the survival of cultured human brain cancer cells U251 MG.

FIGS. 13A and 13B are graphs depicting the in vitro effect of increasing concentrations of SL-11043 (Compound 48, ●) and SL-11044 (Compound 47, ○) on the survival of cultured human brain cancer cells U215 MG.

FIGS. 14A and 14B are graphs depicting the in vitro effect of increasing concentrations of SL-11047 (Compound 58, ●) and SL-11048 (Compound 57, ○) on the survival of cultured human brain cancer cells U251 MG.

FIGS. 15A and 15B are graphs depicting the in vitro effect of increasing concentrations of SL-11037 (Compound 28, ●) and SL-11038 (Compound 23, ○) on the survival of cultured human lung cancer cells A549.

FIGS. 16A and 16B are graphs depicting the in vitro effect of increasing concentrations of SL-11043 (Compound 48, ●) and SL-11044 (Compound 47, ○) on the survival of cultured human lung cancer cells A549.

FIGS. 17A and 17B are graphs depicting the in vitro effect of increasing concentrations of SL-11047 (Compound 58, ●) and SL-11048 (Compound 57, ○) on the survival of cultured human lung cancer cells A549.

FIGS. 18A and 18B are graphs depicting the in vitro effect of increasing concentrations of SL-11037 (Compound 28, ●) and SL-11038 (Compound 23, ○) on the survival of cultured human prostate cancer cells PC3.

FIGS. 19A and 19B are graphs depicting the in vitro effect of increasing concentrations of SL-11043 (Compound 48, ●) and SL-11044 (Compound 47, ○) on the survival of cultured human prostate cancer cells PC3.

FIGS. 20A and 20B are graphs depicting the in vitro effect of increasing concentrations of SL-11047 (Compound 58, ●) and SL-11048 (Compound 57, ○) on the survival of cultured human prostate cancer cells PC3.

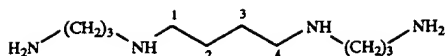
DETAILED DESCRIPTION OF THE INVENTION

Reference is made throughout the Detailed Description to the various reaction Schemes and Tables included herein. For sake of clarity and brevity, reference numerals have been assigned to each unique chemical structure described. These reference numerals are used consistently throughout the disclosure to unambiguously designate the chemical entities discussed.

1. Conformationally Restricted Polyamines: Synthetic Approach

The manufacture of bioactive spermine ligands which may affect the structure of chromatin can be illustrated by the introduction of cyclopropyl and cyclobutyl constraints into the flexible spermine molecule. Spermine appears as follows:

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The first targeted location was the central 1,4-diaminobutane segment. In its staggered conformation, four semi-eclipsed conformational rotamers are possible around the diaminobutane segment. The four have enantiomeric relationships. Introduction of a bond between the C-1 and C-3 positions or the C-2 and C-4 positions of the central diaminobutane segment generates a cyclopropane ring. Introduction of an additional bond between the C-2 and C-3 positions generates a conformationally restricted alkene derivative. Cyclobutyl, cyclopentyl, and cyclohexyl moieties can be introduced into the structure following the same strategy.

Using this approach four conformationally semi-rigid structures were obtained which mimic the four semi-eclipsed conformational structures of spermine. Two of the semi-rigid structures are epimers of the other two.

For purposes of the present invention, it is important to note that the cis and trans isomers of the subject compounds assume very distinct three-dimensional conformations due to the restricted bond rotation afforded by the centrally-located ring structure or unsaturation. All geometric isomers (optically active or otherwise), including pure isolated cis forms and pure isolated trans forms of the subject compounds, and mixtures thereof, are explicitly within the scope of the present invention. Additionally, all positional isomers of the subject compounds are explicitly within the scope of the present invention. When A or D is a cyclical moiety, the two B substituents or the amino moieties, respectively, may be oriented in the 1,2 or 1,3 or 1,4 position with respect to each other.

(a) Spermine Analogs Containing a Cyclopropyl Ring

Cis and trans cyclopropyl analogs of spermine were prepared via the reactions illustrated in Schemes 1-5A.

With reference to Schemes 1 and 2, the cyclopropyl diesters 1 and 2 were first converted into their hydrazides

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103 and 4, and the hydrazides converted into the diamines 5 and 6, respectively. The diamines 5 and 6 were then mesitylated to give the amides 7 and 8, and the amides were then alkylated with 9 to give 10 and 11, respectively. Hydrolysis of the protective groups yielded the trans analog 12 and the cis analog 13.

Referring now to Scheme 3, in a separate reaction, the trans cyclopropyl diester 1 was converted into the amide 14 by reaction with benzylamine (BnNH₂), the amide reduced to the amine 15, and the amine alkylated to 16. The phthalyl residues were then cleaved with hydrazine to give 17. Compound 17 was then either deprotected by hydrogenolysis to give 18; or fully alkylated to 19, and the benzyl residues cleaved by hydrogenolysis to give 20.

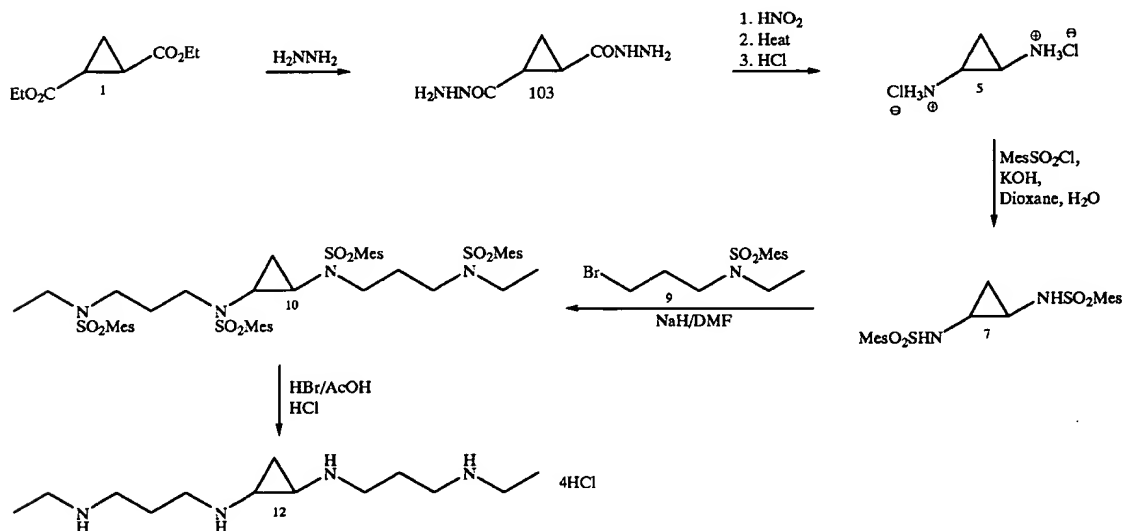
With reference to Scheme 4, the amine 15 was also alkylated with 21 to give 22. Compound 22 was then deprotected to yield the trans cyclopropyl analog 23.

An alternative (and preferred) route to 23 is given in Scheme 4A. Here, 3-ethylamino propionitrile 101 was converted into the corresponding amine 102, which was then mesitylated to yield 3. In a parallel synthesis, the cis diester 1 was reduced to the dialcohol 15', which was then mesitylated to yield the dimesityl derivative 16'. Reacting 3 and 16' in the presence of sodium hydride yields 22'. In the same fashion as Scheme 4, 22' was then deprotected to yield the trans cyclopropyl analog 23.

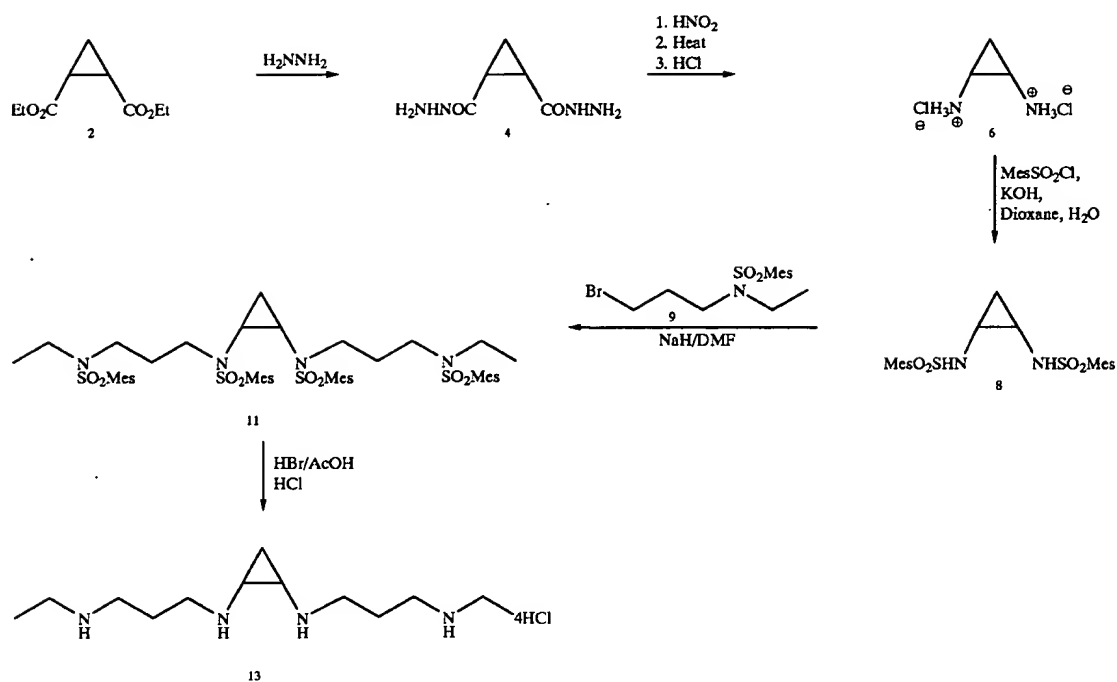
Referring now to Scheme 5, in a separate reaction, the cis cyclopropyl diester 2 was reduced to the dialcohol 24. The dialcohol was then converted into the amine 25, and the amine protected by mesitylation to 26. Compound 26 was then alkylated with 9 to yield 27, and then deprotected to yield the cis cyclopropyl tetramine 28.

An alternative (and preferred) route to 28 is given in Scheme 5A. Here, the cis cyclopropyl diester 2 is reduced to the dialcohol 24 in the same fashion as in Scheme 5. Compound 24 was then protected by mesitylation to yield 25'. Compound 25' was then reacted with 3 to yield 27. Deprotecting yields the tetramine 28.

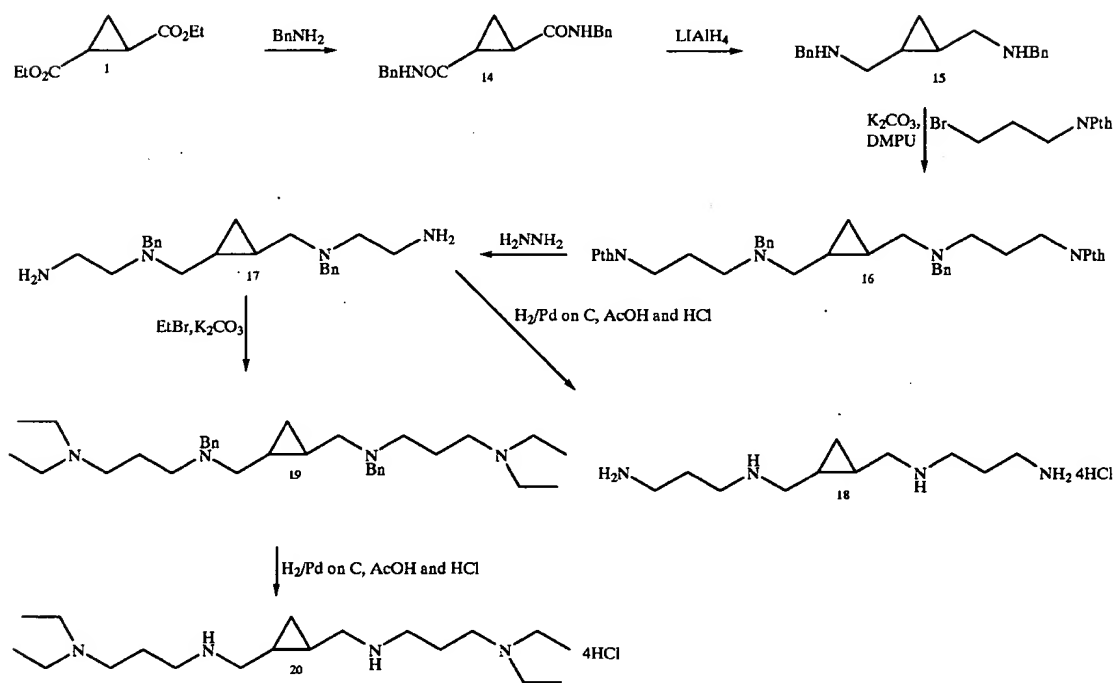
SCHEME 1



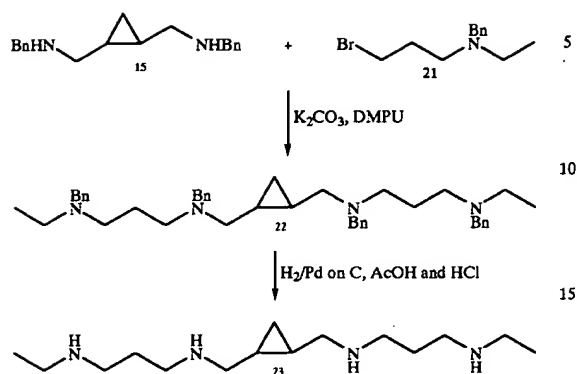
SCHEME 2

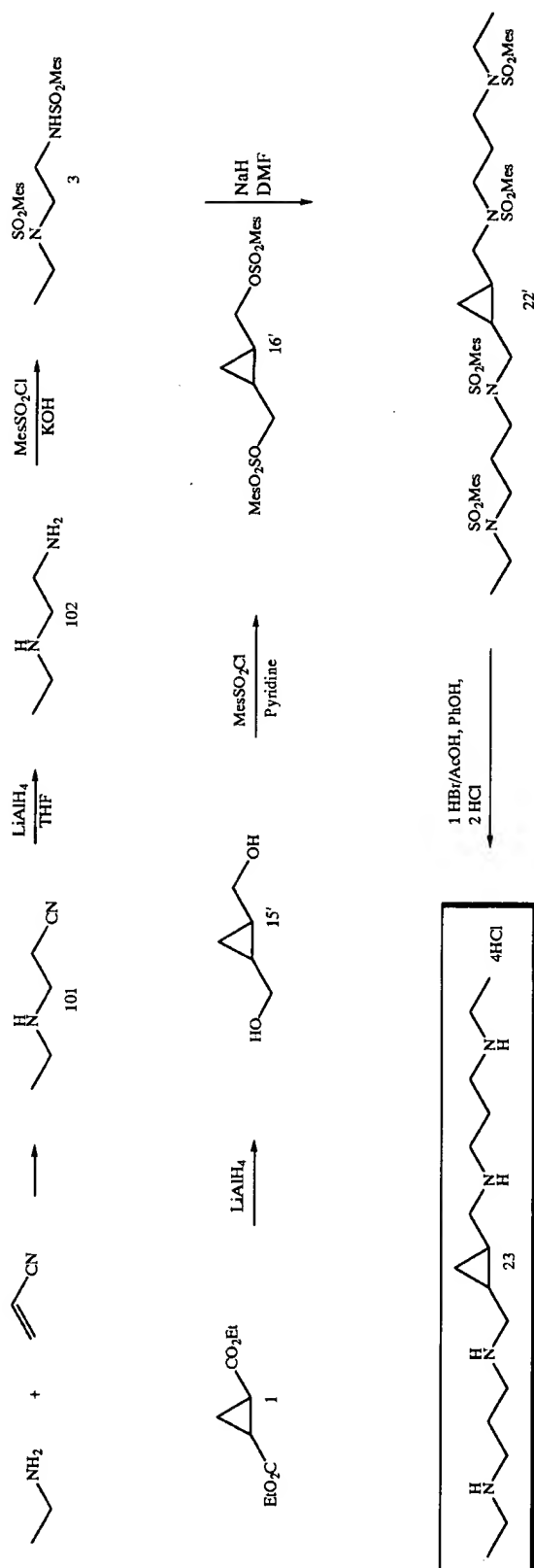


SCHEME 3



SCHEME 4

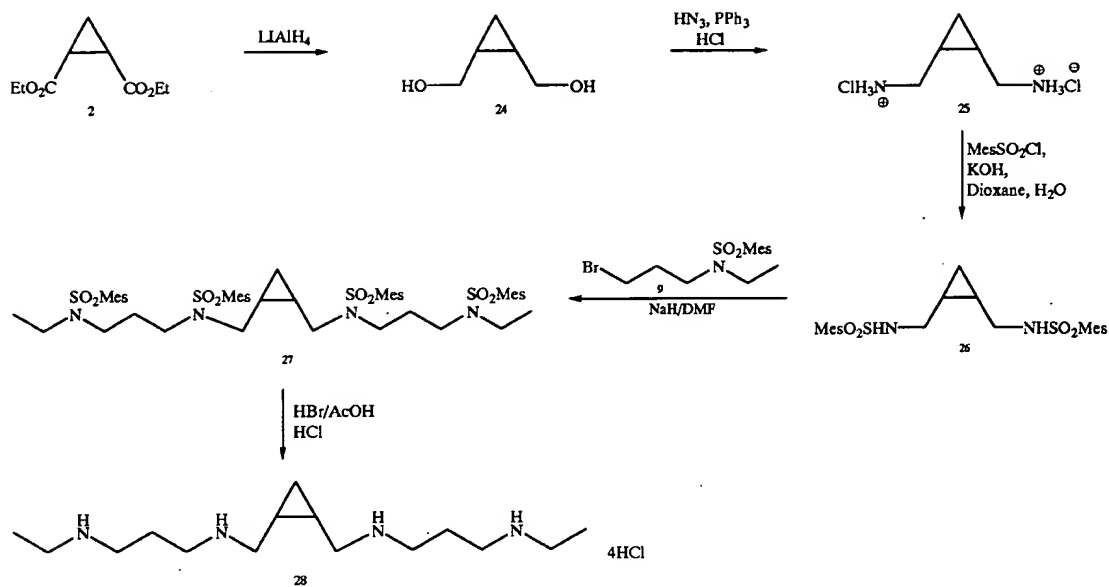




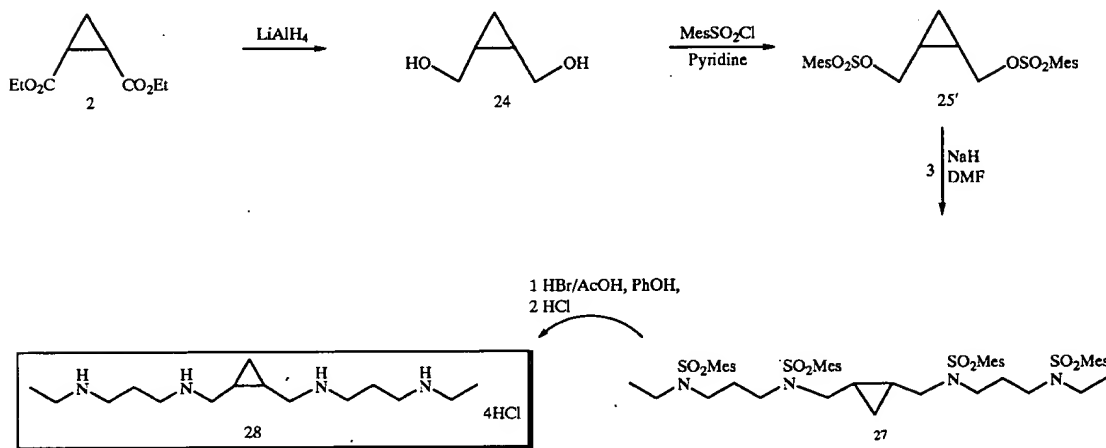
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SCHEME 5



SCHEME 5A



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(b) Spermine Analogs Containing a Cyclobutyl Ring

Cis and trans cyclobutyl analogs of spermine were prepared via the reactions illustrated in Schemes 6-9A.

Referring now to Schemes 6 and 7, the synthesis of the cyclobutyl derivatives started with the trans and cis 1,2-diaminobutanes 29 and 30, respectively. These compounds were first converted to the amides 31 and 32, and then alkylated to give 33 and 34, respectively. Compounds 33 and 34 were then deprotected to yield the trans tetramine 35 (Scheme 6) and the cis tetramine 36 (Scheme 7).

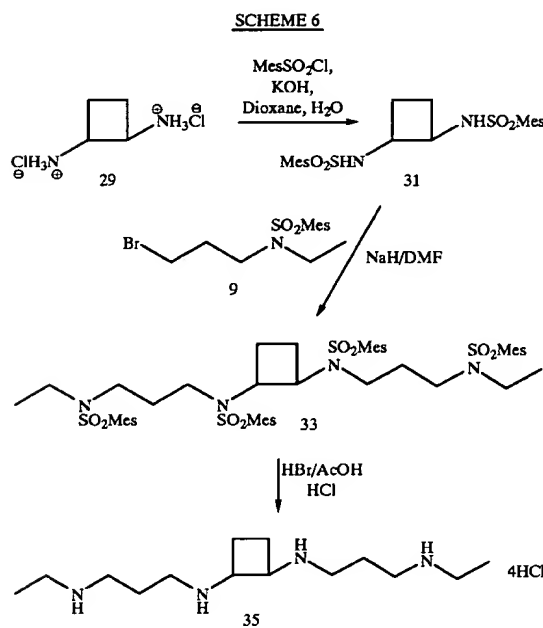
With reference to Schemes 8 and 9, in separate reactions, the trans cyclobutyl diester 37 and the cis cyclobutyl diester

38 were reduced to the respective dialcohols 39 and 40, the dialcohols converted into the diamines 41 and 42. The diamines 41 and 42 were then protected by mesitylation to yield 43 and 44, respectively. These compounds were then alkylated to give 45 and 46. The protecting groups were then removed to yield the trans cyclobutyl tetramine 47 (Scheme 8) and the cis tetramine 48 (Scheme 9).

Alternative (and preferred) routes to 47 and 48 are given in Schemes 8A and 9A, respectively. The cis and trans diesters 37 and 38 were reduced to the respective dialcohols 39 and 40 in the same fashion as in Schemes 8 and 9. Compounds 39 and 40 were then mesitylated to yield 41' and

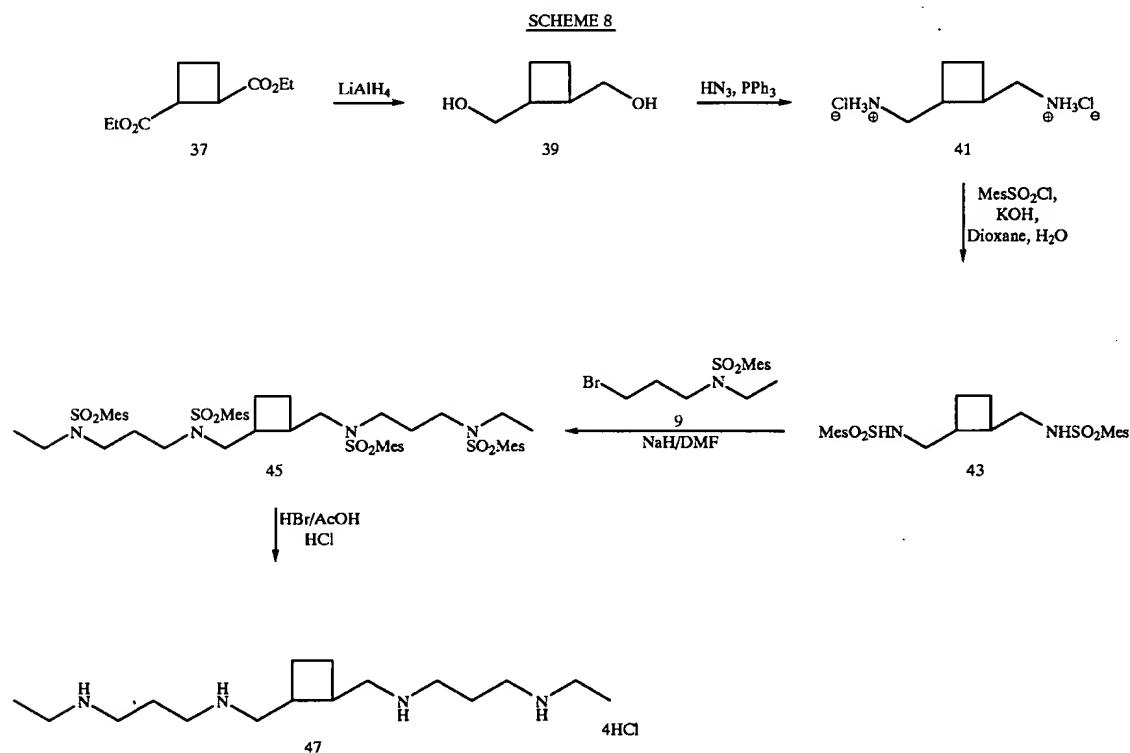
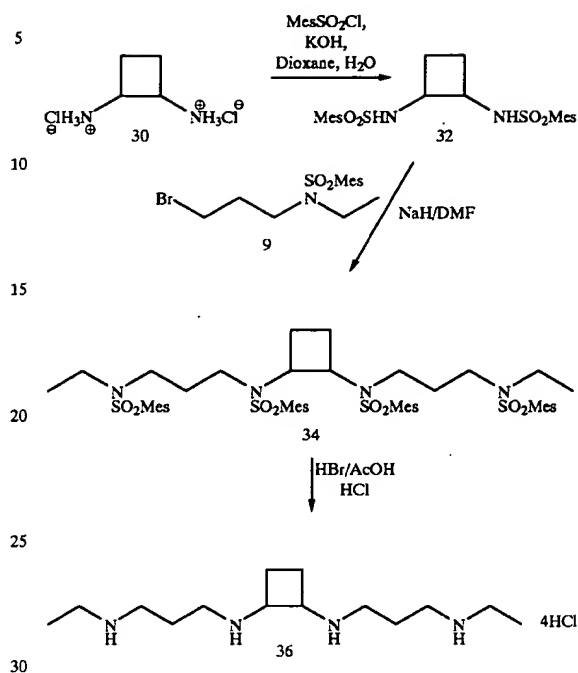
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42', respectively. Reaction of 41' and 42' with 3 yields 45 (Scheme 8A) and 46 (Scheme 9A). Deprotecting yields the desired products 47 and 48.



16

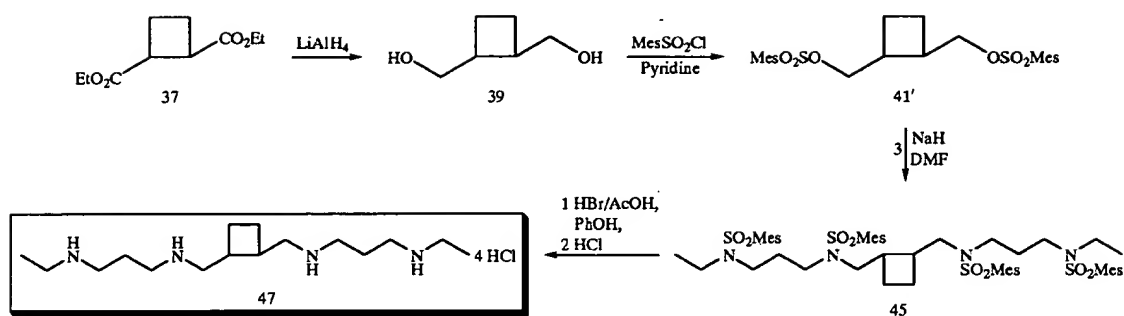
SCHEME 7



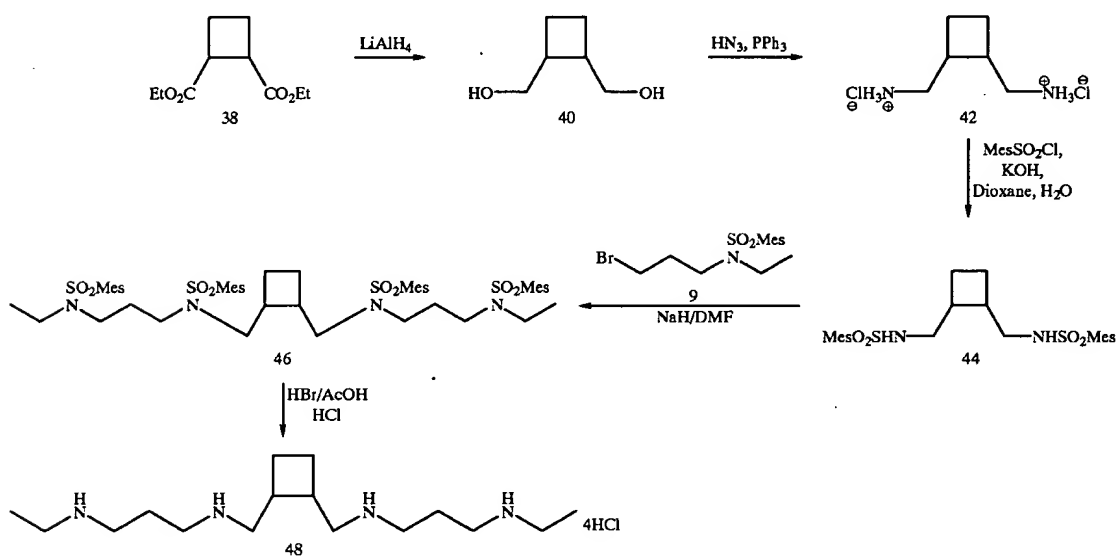
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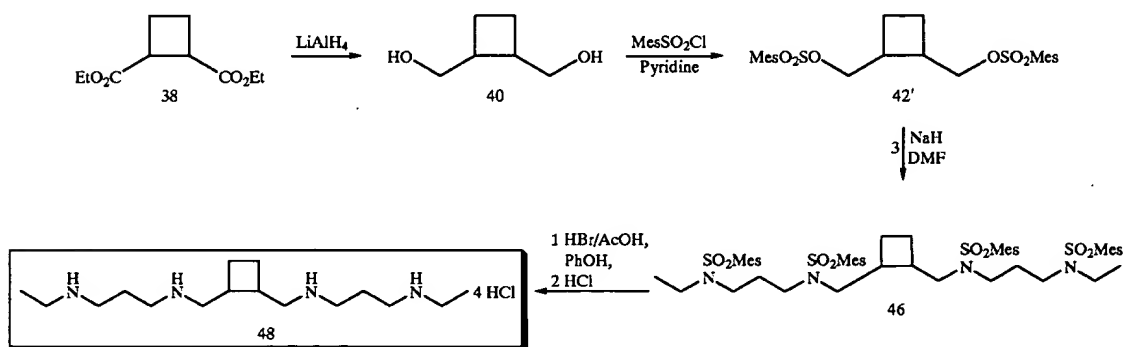
SCHEME 8A



SCHEME 9



SCHEME 9A



19

(c) Spermine Analogs Containing an Unsaturation

Cis and trans unsaturated analogs of spermine were prepared via the reactions illustrated in Schemes 10, 10A, 11, and 11A.

Referring to Scheme 10, the trans diester 49 was reduced to the dialcohol 50, which was then converted into the trans diamine 51. Referring to Scheme 11, the cis diamine 52 was obtained from the commercially available cis dialcohol 43'. With reference to both Scheme 10 and Scheme 11, compounds 51 and 52 were protected by mesitylation to give 53 and 54, respectively. Compounds 53 and 54 were alkylated to 55 and 56, and lastly deprotected to yield the trans tetramine 57 (Scheme 10) and the cis tetramine 58 (Scheme 11).

Alternative (and preferred) routes to 57 and 58 are given in Schemes 10A and 11A, respectively. The cis and trans dialcohols 50' and 50 were obtained in the same fashion as in Schemes 10 and 11. Compounds 50' and 50 were then mesitylated to yield 51' and 52', respectively. Reaction of 51'

20

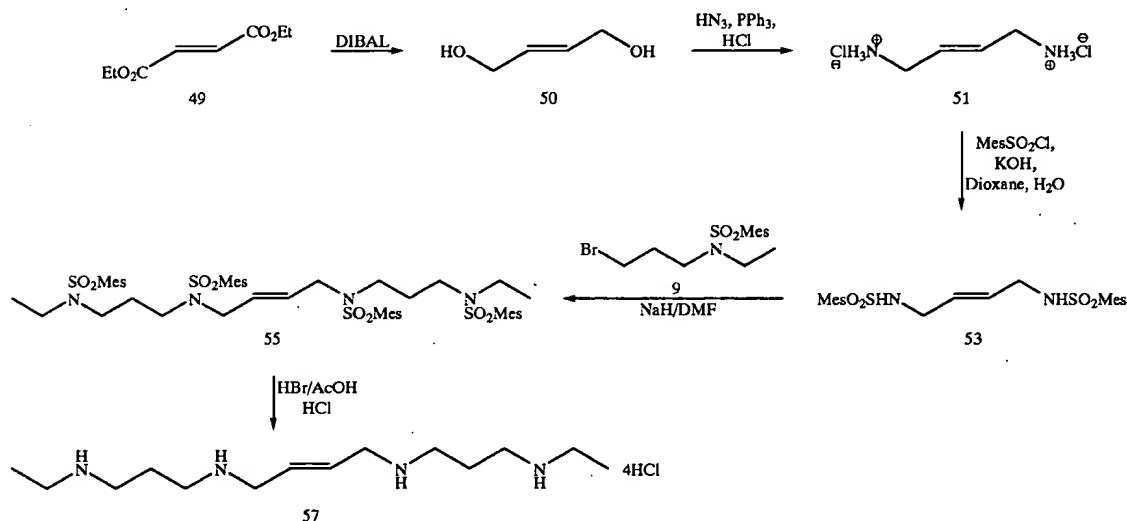
and 52' with 3 yields 55 (Scheme 10A) and 56 (Scheme 11A). Deprotecting yields the desired products 57 and 58.

Following the above general protocols, and using suitable and well known starting reagents, all of the compounds of Formula I, including those where A and D are independently C₅ or C₆ cycloalkyl, cycloalkenyl, or cycloaryl, can be readily obtained. An illustrative listing of compounds of Formula I are presented in Table 1.

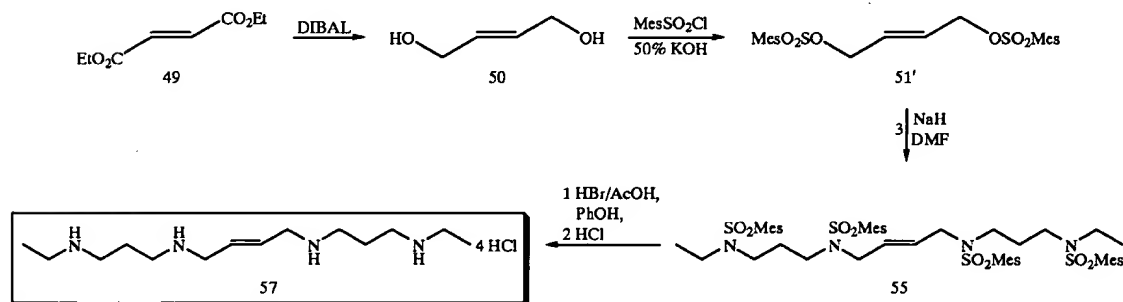
See the Examples, below, for illustrative syntheses of the compounds shown in Table 1.

The pure compounds, as well as pharmaceutically-suitable salts thereof, are explicitly within the scope of the present invention. By the term "pharmaceutically-suitable salts" is meant any salt form of the subject compounds which renders them more amenable to administration by a chosen route. A wide range of such salts are well known to those of skill in the pharmaceutical art. The preferred pharmaceutically-suitable salts are acid addition salts such as chlorides, bromides, iodides and the like.

SCHEME 10



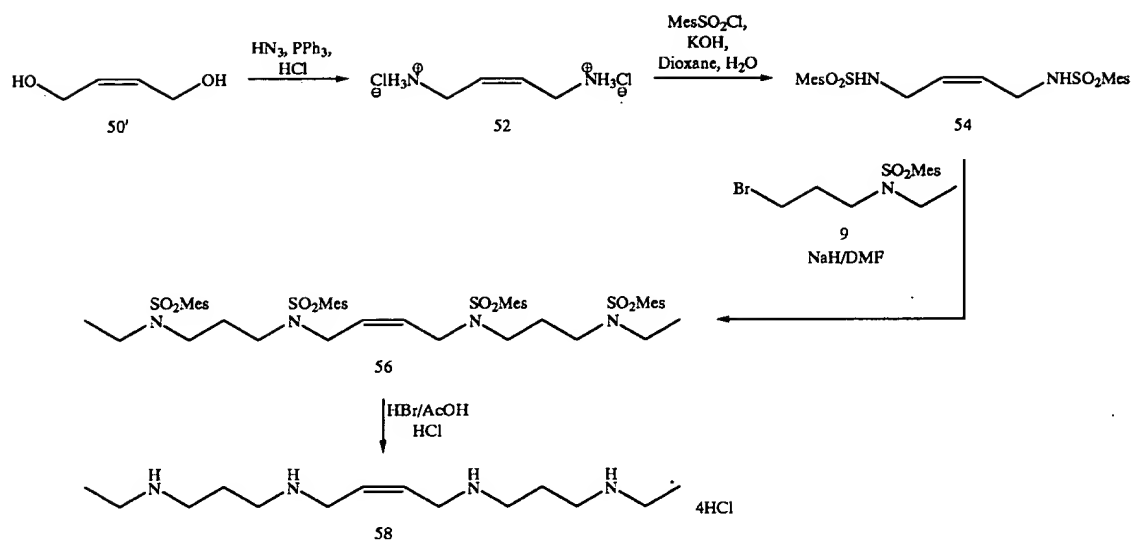
SCHEME 10A



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SCHEME 11



SCHEME 11A

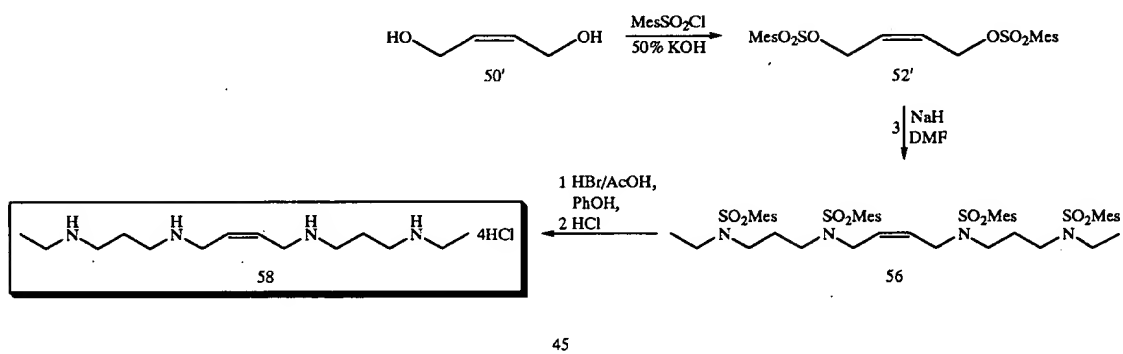
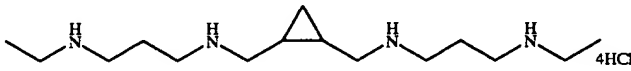
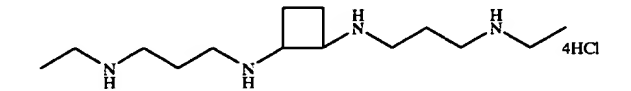
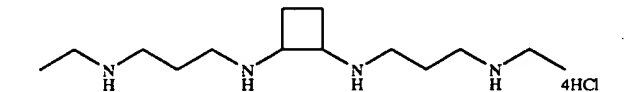
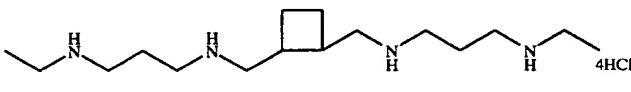
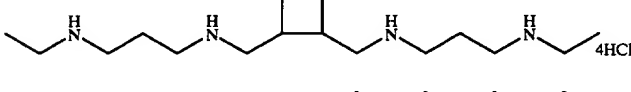
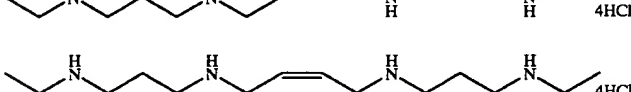
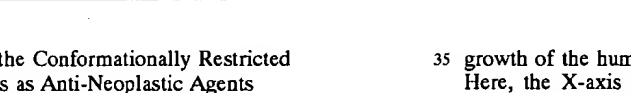


TABLE 1

Compound No.		
12	<chem>CH3CH2NHCH2CH2CH2NHCH2CH2CH2N1CC1NCH2CH2CH2NHCH3</chem> $\cdot 4\text{HCl}$	SL-11027
13	<chem>CH3CH2NHCH2CH2CH2NHCH2CH2CH2N1CC1NCH2CH2CH2NCH2CH2CH2NHCH3</chem> $\cdot 4\text{HCl}$	SL-11033
23	<chem>CH3CH2NHCH2CH2CH2NHCH2CH2CH2N1CC1NCH2CH2CH2NCH2CH2CH2NCH2CH2CH2NHCH3</chem> $\cdot 4\text{HCl}$	SL-11038

TABLE 1-continued

Compound No.		
28		SL-11037
35		SL-11028
36		SL-11034
47		LS-11044
48		SL-11043
57		SL-11048
58		SL-11047

2. Utility of the Conformationally Restricted Polyamines as Anti-Neoplastic Agents

To assess the utility of the subject compounds in the treatment of neoplastic cell growth, the ability of the compounds to inhibit the in vitro growth of several commonly used cancer models was studied. The subject polyamines induce cell death in several neoplastic cell lines at drug concentrations smaller than 10 μ M. In serial dilution, the restricted conformation polyamines of the present invention have been shown to inhibit cell growth and/or cause cell death in accepted in vitro test cultures for human breast cancer (MCF7), brain cancer (U251MG NCI), lung cancer (A549), colon cancer (HT29), and prostate cancer (PC3) at minute concentrations heretofore undescribed in the scientific literature.

The drawing figures are graphs depicting the results of a series of experiments which illustrate the ability of the subject polyamines to induce cell death in different neoplastic cell lines. These figures are described in detail in the Examples Section, below.

Referring to FIGS. 1-6, each of these graphs has as its X-axis the concentration of the particular compound being tested. The Y-axis of the graphs depicted in FIGS. 1-6 is a linear scale representing the fraction of cell survival in each of the cultures tested. Here, the neoplastic cell line MCF7 was used. This cell line is a human breast cancer cell line. These figures clearly illustrate the utility of the subject polyamines to inhibit the growth of human breast cancer. The 6 compounds whose in vitro activity is depicted in FIGS. 1-6 display ED₅₀ values ranging from 1.34 to 1.79 μ M.

FIGS. 7 and 8 depict the effect of fixed doses of 10 μ M and 40 μ M of the subject compounds, respectively, on the

growth of the human brain cancer cell line U251MG-NCI. Here, the X-axis represents time in days, and the Y-axis represents the total number of viable cells. FIGS. 7 and 8 further demonstrate the utility of subject compounds to inhibit neoplastic cell growth.

FIGS. 9A, 9B, 10A, 10B, 11A, and 11B show the in vitro effect of several of the subject compounds on HT29 cells. FIGS. 12A, 12B, 13A, 13B, 14A, and 14B show the in vitro effect of several of the subject compounds on U251 MG cells. FIGS. 15A, 15B, 16A, 16B, 17A, and 17B show the in vitro effect of several of the subject compounds on A 549 cells. Lastly, FIGS. 18A, 18B, 19A, 19B, 20A, and 21B show the in vitro effect of several of the subject compounds on PC 3 cells.

3. Administration and Pharmaceutical Unit Dosage Forms

The above-described compounds being effective to inhibit the growth of cancer cells, the compounds are suitable for the therapeutic treatment of neoplastic conditions in mammals, including humans. Cancer cell growth inhibition at pharmacologically-acceptable concentrations has been shown in human breast cancer, brain cancer, lung cancer, colon cancer, and prostate cancer cell lines.

Administration of the subject conformationally restricted polyamines to a human or non-human patient can be accomplished by any means known. The preferred administration route is parenteral, including intravenous administration, intraarterial administration, intratumor administration, intramuscular administration, intraperitoneal administration, and subcutaneous administration in combination with a pharmaceutical carrier suitable for the chosen administration route. The treatment method is also amenable to oral administration.

It must be noted, as with all pharmaceuticals, the concentration or amount of the polyamine administered will

vary depending upon the severity of the ailment being treated, the mode of administration, the condition and age of the subject being treated, and the particular polyamine or combination of polyamines being used.

The compounds described herein are administratable in the form of tablets, pills, powder mixtures, capsules, injectables, solutions, suppositories, emulsions, dispersions, food premixes, and in other suitable forms. The pharmaceutical dosage form which contains the compounds described herein is conveniently admixed with a non-toxic pharmaceutical organic carrier or a non-toxic pharmaceutical inorganic carrier. Typical pharmaceutically-acceptable carriers include, for example, mannitol, urea, dextrans, lactose, potato and maize starches, magnesium stearate, talc, vegetable oils, polyalkylene glycols, ethyl cellulose, poly(vinylpyrrolidone), calcium carbonate, ethyl oleate, isopropyl myristate, benzyl benzoate, sodium carbonate, gelatin, potassium carbonate, silicic acid, and other conventionally employed acceptable carriers. The pharmaceutical dosage form may also contain non-toxic auxiliary substances such as emulsifying, preserving, or wetting agents, and the like.

Solid forms, such as tablets, capsules and powders, can be fabricated using conventional tableting and capsule-filling machinery, which is well known in the art. Solid dosage forms may contain any number of additional non-active ingredients known to the art, including excipients, lubricants, dessicants, binders, colorants, disintegrating agents, dry flow modifiers, preservatives, and the like.

Liquid forms for ingestion can be formulated using known liquid carriers, including aqueous and non-aqueous carriers, suspensions, oil-in-water and/or water-in-oil emulsions, and the like. Liquid formulation may also contain any number of additional non-active ingredients, including colorants, fragrance, flavorings, viscosity modifiers, preservatives, stabilizers, and the like.

For parenteral administration, the subject compounds may be administered as injectable dosages of a solution or suspension of the compound in a physiologically-acceptable diluent or sterile liquid carrier such as water or oil, with or without additional surfactants or adjuvants. An illustrative list of carrier oils would include animal and vegetable oils (peanut oil, soy bean oil), petroleum-derived oils (mineral oil), and synthetic oils. In general, for injectable unit doses, water, saline, aqueous dextrose and related sugar solutions, and ethanol and glycol solutions such as propylene glycol or polyethylene glycol are preferred liquid carriers.

The pharmaceutical unit dosage chosen is preferably fabricated and administered to provide a concentration of drug at the point of contact with the cancer cell of from 1 μ M to 10 mM. More preferred is a concentration of from 1 to 100 μ M. This concentration will, of course, depend on the chosen route of administration and the mass of the subject being treated.

EXAMPLES

Chemical Synthesis Examples

The following Examples are an illustrative, not exclusive, listing of the manufacture of several compounds according to the present invention. The Examples are included herein solely to aid in a more complete understanding of the present invention. The Examples do not limit the invention disclosed and claimed herein in any fashion. Reference numerals are to the Reaction Schemes described above.

All melting points are uncorrected. NMR spectra were recorded on 300 MHz spectrometer with TMS as internal standard. Purification of the products was carried out using

silica gel 60 (230–400 mesh) and the appropriate solvent system. HPLCs were run on a Waters 8 \times 10 "NOVAPAK" cartridge (Millipore Corporation, Marlboro, Mass.) using reverse phase and a gradient of acetonitrile and sodium acetate buffer, pH 4.5.

Compounds 1 and 2 were prepared according to the method of Ashton et al. (1988).

Compound 1: ^1H NMR (CDCl_3): δ 1.26 (t, J=7 Hz, 6H, CH_2CH_3), 1.43 (dd, J=7.0, 8.0 Hz, 2H, CyPrCH₂), 2.16 (dd, J=7.0, 8.0 Hz, 2H, CyPrCH), 4.16 (q, J=7 Hz, 4 H, CH_2CH_3). ^{13}C -NMR (CDCl_3): 14.12, 15.25, 22.31, 61.01, 171.75.

Compound 2: ^1H NMR (CDCl_3): δ 1.25 (m, 1H, CyPrCH₂), 1.26 (t, J=7 Hz, 6H, CH_2CH_3), 1.70 (ddd, J=5.0, 6.0, 5.5 Hz, 1H, CyPrCH₂), 2.06 (dd, J=6.0, 9.0 Hz, 2H, CyPrCH), 4.17 (q, J=7 Hz, 4H, CH_2CH_3).

Compound 103: To a solution of ethyl ester 1 (3.5 mL, 20 mmol) in ethanol (10 mL) was added hydrazine monohydrate (3.38 mL, 80 mmol). The mixture was heated to reflux overnight to afford 103 as a white solid. The reaction was cooled to room temperature (RT), diluted with 20 mL CHCl_3 , and the solid was filtered. Yield=80%; m.p. 215–217° C.; ^1H NMR (D_2O): δ 1.35 (dd, J=7.0, 8.0 Hz, 2H, CyPrCH), 2.04 (dd, J=7.0, 8.0 Hz, 1H, CyPrCH); ^{13}C -NMR (D_2O): 15.5, 24, 176.

Compound 4: This compound was synthesized via the route described above for compound 103. Yield was 87%; m.p. 196–198° C.; ^1H NMR (D_2O): δ 1.25 (m, 1H, CyPrCH₂), 1.45 (ddd, J=5.0, 6.0, 5.5 Hz, 1H, CyPrCH₂), 2.05 (dd, J=6.0, 9.6 Hz, 2H, CyPrCH); ^{13}C -NMR (D_2O): 12.97, 23.88, 175.0.

Compound 5: Concentrated HCl was placed into a 125 mL conical flask and 9 g of crushed ice added thereto. Compound 103 (1.58 g, 10 mmol) was dissolved in this solution and then ethyl ether (10 mL) was added. While maintaining the temperature below 10° C., a NaNO_2 solution (1.73 g, 25 mmol in 4 mL H_2O) was added slowly. The organic layer was separated, the aqueous layer extracted with ether (3 \times 20 mL), and the organic fractions combined and dried over CaCl_2 . The ether solution was filtered into a 250 mL round bottom flask and diluted with anhydrous toluene (30 mL). Ether was distilled-off using a fractionating column condenser. The remaining toluene solution was heated to 85° C. until the evolution of nitrogen ceased. Stirring at 85° C. was continued for and additional 10 min. While still hot, this solution was poured into preheated (60° C.) concentrated HCl (8 mL). Toluene was distilled off under vacuum and anhydrous ethanol (15 mL) was added to the flask and then distilled-off. This process was repeated twice to afford a cream-colored solid, which was digested with ethanol and suction-filtered to afford 0.645 g (45%) of pure 5; m.p. 220° C. (dec); ^1H NMR (D_2O) δ 1.53 (t, J=6.0 Hz, 2H), 3.2 (t, J=6.0 Hz, 2H).

Compound 6: This compound was synthesized in the same manner as 5. Yield was 58%; m.p. 220° C.; ^1H NMR (D_2O) δ 1.25 (m, 2H, CyPrCH₂), 2.9 (dd, 2H, J=6.0, 8.0 Hz, CyPrCH).

Compound 7: Compound 5 (145 mg, 1 mmol) was dissolved in 4 mL of dioxane/water (1:1), while maintaining a pH ca. 11 by the addition of 5% KOH. Mesitylenesulfonyl chloride (875 mg, 4 mmol) in 5 mL dioxane was then added slowly. The upper layer of the mixture was carefully decanted and the gummy residue was triturated with hexanes to afford 330 mg (76%) of 7 as a white solid, which was recrystallized from CHCl_3 /hexanes. m.p. 189–191° C.; ^1H NMR (CDCl_3): δ 0.9 (t, J=8.0 Hz, 2H CyPrCH₂), 2.24 (t,

J=8 Hz, 2H, CyPrCH), 2.30 (s, 6H, 2 CH₃), 2.55 (s, 12H, 4 CH₃), 5.00 (s, 2H, NH), 6.75 (s, 4H, aromatic); ¹³C-NMR (CDCl₃): 14.16, 20.98, 22.92, 30.92, 132.11, 132.89, 139.20, 142.79; MS (m/z): 436.53 (M⁺), 421, 253, 183, 171, 119, 91; Anal: C₂₁H₂₈N₂S₂O₄.

Compound 8: This compound was synthesized in the same manner as 7. Yield was 95%; mp 175–177° C.; ¹H-NMR (CDCl₃): 80.80 (m, 1H, CyPrCH₂), 0.95 (m, 1H, CyPrCH₂), 2.25 (m, 2H, CyPrCH), 2.34 (s, 6H, 2 CH₃), 2.65 (s, 12H, 4 CH₃), 5.25 (br, 2H, NH), 7.00 (s, 4H, aromatic); ¹³C-NMR (CDCl₃): 13.5, 20.96, 22.96, 28.00, 132.08, 133.00, 139.45, 142.09.

Compound 10: This compound was synthesized by the procedure of Bergeron et al. (1994). NaH (95%, 0.111 g, 4.4 mmol) was added to 7 (0.872 g, 2 mmol) in anhydrous dimethylformamide (40 mL) at 0° C. The mixture was stirred for 30 min at 0° C. Compound 9 (1.531 g, 4.4 mmol) in anhydrous DMF (50 mL) was added slowly, stirred for another 15 min. at 0° C. and then stirred overnight at room temperature. The reaction was quenched with water (8 mL), followed by extraction with ether (3x25 mL). The combined organic layers were washed with water (4x30 mL) and brine (2x25 mL). The solvents were removed under vacuum to afford a gum, which was filtered on a silica gel column using 8:1 hexanes:ethylacetate as an eluant. Obtained was 1.7 g (77%) of 10 as a white solid. m.p. 60–6° C.; ¹H-NMR (CDCl₃) 80.57 (dd, J=6 and 8 Hz, 2H, CyPrCH₂), 0.99 (t, J=8 Hz, 6 H, 2 CH₃), 1.7–1.9 (m, 4H, NCH₂CH₂), 2.25 (s, 6H, 2 CH₃), 2.29 (s, 6H, 2 CH₃), 2.50 (s, 12H, 4 CH₃), 2.55 (s, 12 H, 4 CH₃), 2.56–2.67 (m, 2H, CyPrCH), 2.87–3.19 (m, 12H, 6 NCH₂), 6.86 (s, 4H, aromatic), 6.90 (s, 4H, aromatic); MS (m/z) 971.4 (M⁺), 787.3, 605.3, 295.1, 183, 119.1 (100%); Anal: (C₄₉H₇₀N₄S₄O₈).

Compound 11: This compound was obtained (29%) from 7 as described above for compound 10. mp=74–75° C.; ¹H-NMR (CDCl₃) 80.25–0.36 (m, 1H, CyPrCH₂), 0.99 (t, J=8 Hz, 6H, 2CH₃), 1.00–1.11 (m, 1H, CyPrCH₂), 1.50–1.70 (m, 4H, NCH₂CH₂), 2.30 (s, 12H, 4CH₃), 2.55 (s, 12H, 4CH₃), 2.58 (s, 12H, 4CH₃), 2.56–2.70 (m, 2H, CyPrCH), 2.90–3.20 (m, 12H, 6NCH₂), 6.90 (s, 4H, aromatic), 6.93 (s, 4H, aromatic). MS (m/z) 971.4 (M⁺), 787.3, 605.3, 154 (100%), 119. Anal (C₄₉H₇₀N₄S₄O₈).

Compound 12: This compound was synthesized by the procedure of Bergeron et al. (1994) in 77% yield. Recrystallized from EtOH, mp=240° C. (dec.); ¹H-NMR (D₂O/Acetone-d₆) 81.23 (t, J=6.8, 2H, CyPrCH₂), 1.31 (t, J=7.3 Hz, 6H, 3CH₃), 2.00–2.10 (m, 4H, NCCH₂), 2.81 (t, J=6.6 Hz, 2H, CyPrCH), 3.00–3.20 (m, 12H, 6 NCH₂), ¹³C-NMR (D₂O, Acetone-d₆) 813.33, 13.73, 26.64, 37.63, 45.76, 47.27, 47.83; MS (m/z) 243.4 (M⁺, 100%), 163.3, 141.3.

Compound 13: This compound was obtained from 11 in 28% yield, mp=240° C. (dec). ¹H-NMR (D₂O/Acetone-d₆) 80.72–0.81 (m, 1H, CyPrCH₂), 1.16 (ddd, J=7.2, 8.1, 7.2 Hz, 1H, CyPrCH₂), 1.29 (t, J=7.4 Hz, 6H, 2CH₃), 2.04 (pent, J=7.5 Hz, 4H, NCCH₂), 2.69 (dd, J=6.8, 7.4 Hz, 2H, CyPrCH), 2.98–3.24 (m, 12H, NCH₂); ¹³C-NMR (D₂O, Acetone-d₆) 812.44, 13.43, 27.12, 35.86, 45.89, 47.48, 48.06; MS (m/z): 243.3 (M⁺, 100%), 163.2, 142.7.

Compound 14: Compound 1 (3.5 mL, 20 mmol) was dissolved in toluene (30 mL) and benzylamine (4.61 mL, 44 mmol) was added to the solution. The mixture was stirred overnight at 80° C. to afford 14 as white solid which was filtered and washed with EtOH, yield 85%; mp=237–238° C.

Compound 15: LiAlH₄ (200 mg, 5.26 mmol) was suspended in anhydrous THF (10 mL) and kept under argon. Compound 14 (500 mg, 1.62 mmol) was added to the

suspension in portions. Though 14 was insoluble at RT, it was slowly dissolved while refluxing overnight. The reaction was quenched with MeOH and the solvents were evaporated under vacuum to dryness. The residue was then extracted with CHCl₃ (2x15 mL). The organic layer was cooled on an ice-salt mixture to precipitate by-products which were filtered off. CHCl₃ was then removed under vacuum to afford 15. It was purified by column chromatography using silica gel and 5:1 hexanes:ethylacetate as an eluant. This gave 45% of 15 as a thick gum. ¹H-NMR (CDCl₃) 80.36 (dd, J=7.7, 7.2 Hz, 2H, CyPrCH₂), 0.84 (t, J=7.7 Hz, 2H, CyPrCH), 1.91 (br, 2H, NH), 2.44–2.60 (m, 4H, 2NCH₂), 3.81 (s, 4H, Benzyl), 7.30 (s, 10H, aromatic); ¹³C-NMR (CDCl₃) 815.27, 39.12, 53.46, 126.93, 128.20, 128.41, 140.42.

Compound 16: Compound 15 (4.5 g, 16.1 mmol) was dissolved in dimethylpropyleneurea (DMPU) (35 mL), and K₂CO₃ (8.95 g, 64.8 mmol) was added, followed by 3-bromopropylphthalimide (8.67 g, 32.3 mmol) in DMPU (15 mL) and the solution was stirred at 110° C. overnight. The solution was then cooled to 0° C., and the precipitated solids were dissolved in H₂O (50 mL). The mixture was extracted with ether (3x50 mL), the organic layers were washed with water (3x25 mL), dried (MgSO₄), and evaporated in vacuo to afford a thick gum. This gum was subjected to column chromatography on silica gel column using hexanes:ethylacetate (7:3) as an eluant to give 6.1 g (58%) of 16. ¹H-NMR (CDCl₃) 80.31 (dd, J=6.4, 6.8 Hz, 2H, CyPrCH₂), 0.66 (t, J=5.8 Hz, 2H, CyPrCH), 1.83 (pent, J=7.0 Hz, 4H, NCCH₂), 2.35 (dq_{AB}, J=6.3, 13.2 Hz, 4H, NCH₂), 2.56 (t, J=7.0 Hz, 4H, 2NCH₂), 3.60 (s, 4H, Benzyl), 3.69 (t, J=7.4 Hz, 4H, NCH₂), 7.1–7.4 (m, 10H, aromatic), 7.65–7.85 (m, 8H, aromatic); ¹³C-NMR (CDCl₃) 811.10, 15.24, 26.01, 36.43, 50.82, 57.48, 58.37, 123.10, 126.67, 128.10, 128.77, 132.22, 133.78, 139.80, 168.36.

Compound 17: Compound 16 (5.86 g, 8.96 mmol) was dissolved in methanol (100 mL) and hydrazine monohydrate (3.6 mmol) was added and the mixture was heated to reflux overnight. The methanol was evaporated under vacuum, and NH₄OH (4M, 100 mL) added to the residue. This solution was extracted vigorously with CHCl₃ (4x50 mL), the combined organic layers dried (MgSO₄), and CHCl₃ evaporated to afford essentially pure 17 as a thick gum (3.3 g, 94% yield); ¹H-NMR (CDCl₃) 80.34 (dd, J=6.4, 6.8 Hz, 2H, CyPrCH₂), 0.68 (t, J=5.8 Hz, 2H, CyPrCH), 1.62 (pent, J=6.8 Hz, 4H, NCCH₂), 1.70 (br, 4H, NH₂), 2.35 (dq_{AB}, J=6.1, 13.2 Hz, 4H, NCH₂), 2.51 (t, J=6.8 Hz, 4H, NCH₂), 2.69 (t, J=6.7 Hz, 4H, NCH₂), 3.60 (s, 4H, Benzyl), 7.2–7.39 (m, 10H, aromatic); ¹³C-NMR (CDCl₃) 811.09, 15.54, 30.72, 40.49, 51.18, 57.77, 58.62, 126.75, 128.16, 128.76, 140.05.

Compound 18: Compound 17 (0.5 g, 1.27 mmol) was dissolved in acetic acid (10 mL). To this solution was added 10% Pd on activated carbon (200 mg) and the mixture was hydrogenated overnight at 60° C. The catalyst was filtered off and 1M HCl in ether (10 mL) was added to the solution to afford a white solid, which was filtered and recrystallized from MeOH/ether to give 0.210 g (46%) of 18. mp=270° C. (dec); ¹H-NMR ((D₂O) 80.84 (dd, J=6.7, 7.1 Hz, 2H, CyPrCH₂), 1.20 (t, J=6.5 Hz, 2H, CyPrCH), 2.00–2.20 (m, 4H, NCCH₂), 2.05–2.90 (m, 2H, NCH₂), 3.00–3.25 (m, 10H, NCH₂); MS (m/z) 215.1 (M⁺, 100%), 144.9, 132, 124, 98.

Compound 19: Compound 17 (1.5 g, 3.8 mmol) was dissolved in DMPU (10 mL) and K₂CO₃ (1.52 g, 10.98 mmol) added to the solution, followed by the addition of ethylbromide (0.56 mL, 7.6 mmol). The flask was fitted with

a dry-ice reflux condenser, then stirred overnight at 45° C. The mixture was then cooled to 0° C., and the precipitated solids were dissolved in H₂O (100 mL), extracted with ether (3×100 mL), the combined organic layers washed with H₂O (3×50 mL), dried (MgSO₄), and the solvent evaporated to afford a thick gum. The gum was purified by column chromatography over silica gel using ethyl acetate as an eluant (95% yield). ¹H NMR (CDCl₃) δ 0.33 (dd, J=6.5, 7.0 Hz, 2H, CyPrCH₂), 0.65 (t, J=6.5 Hz, 2H, CyPrCH), 1.00 (t, J=7.2 Hz, 12H, 4CH₃), 1.55–1.69 (m, 4H, NCCH₂), 2.25 (dd, J=6.3, 13.2 Hz, 2H, NCH₂), 2.30–2.55 (m, 18H, NCH₂), 3.62 (s, 4H, benzyl), 7.19–7.35 (m, 10H, aromatic).

Compound 20: This compound was obtained as described above for 18 starting from 19 (80% yield). mp=230° C. (dec); ¹H NMR (D₂O) δ 0.90 (t, J=6.8 Hz, 2H, CyPrCH₂), 1.22 (t, J=6.5 Hz, 2H, CyPrCH), 1.33 (t, J=7.3 Hz, 12H, 4CH₃), 3.15–3.35 (m, 4H, NCCH₂), 2.95 (dd, J=8, 13 Hz, NCH₂), 3.15–3.35 (m, 18H, NCH₂); ¹³C-NMR (D₂O) δ 11.83, 13.66, 17.30, 24.35, 47.77, 51.18, 52.05, 54.46; MS (m/z) 327 (M⁺, 100%).

Compound 22: Compound 22 was prepared as described above for 16, using 15 as the reactant (51% yield). ¹H NMR (CDCl₃) δ 0.65 (t, J=6.8 Hz, 2H, CyPrCH₂), 0.90–1.1 (m, 8H, 2CH₃, CyPrCH), 1.50–1.75 (m, 4H, NCCH₂), 2.25–2.55 (m, 16H, NCH₂), 3.49 (s, 8H, benzylic), 7.10–7.40 (m, 20H, aromatic).

Compound 23 (Scheme 4): Compound 23 was prepared (20%) following the procedure used for 18. The product was recrystallized from ethanol/ether; mp=270° C. (dec) ¹H NMR (D₂O) δ 0.88 (dd, J=6.8, 7.0 Hz, 2H, CyPrCH₂), 1.25 (t, J=6.1 Hz, 2H, CyPrCH), 1.30 (t, J=7.3 Hz, 6H, 2CH₃), 2.12 (pent., J=7.9 Hz, 4H, NCCH₂), 2.89 (dd, J=8.5, 12.2 Hz, 2H, NCH₂), 3.08–3.30 (m, 14H, NCH₂); ¹³C-NMR (D₂O) δ 13.62, 14.10, 17.25, 26.35, 46.70, 47.49, 47.81, 54.41; MS (m/z) 271.4 (M⁺H⁺), 307.3 (M⁺H⁺+2 HCl), 156.7, 136.3 (100%).

Compound 23 (via Scheme 4A): Referring specifically to Scheme 4A, 3-ethylamino propionitrile, 101, was prepared by the method of Israel et al. (1964). ¹H-NMR (CDCl₃): δ 1.13 (t, J=7.2 Hz, 3H, CH₃), 2.53 (t, J=6.8 Hz, 2H, NCH₂), 2.69 (q, J=7.2 Hz, 2H, NCH₂), 2.94 (t, J=6.8 Hz, 2H, CNCH₂); ¹³C NMR (CDCl₃): δ 15.09, 18.62, 43.37, 44.82, 118.75.

N,1-Ethylpropane-1,3-diamine, 102: LiAlH₄ (2.54 g, 66.93 mmol) was taken into a flame dried, 3-necked flask. THF (75 ml) was added slowly and stirred at room temperature for 20 min. under argon atmosphere. Compound 101 (4.322 g, 44.1 mmol) in 25 ml THF was added dropwise, stirred its contents over night at 80° C. The reaction mixture was cooled to room temperature and quenched with 150 ml 30% NaOH solution. The organic layer was separated, followed by extraction of aqueous layer with chloroform (100 ml). Combined organic layers were dried over Na₂SO₄. Solvents were evaporated under vacuum to afford 3.28 g (73%) of compound 102 as an oil which was relatively pure from NMR spectrum and used in the next step without further purification. ¹H NMR (CDCl₃): δ 1.11 (t, J=7.2 Hz, 3H, CH₃), 1.64 (pent, J=7.0 Hz, 2H, NCCH₂), 2.55–2.71 (m, 4H, NCH₂), 2.77 (t, J=6.8 Hz, 2H, NCH₂); ¹³C NMR (CDCl₃): δ 15.26, 25.55, 33.89, 40.33, 44.13, 47.60.

N-Ethyl-N-(3-Mesitylene Sulfonylamino Propyl)-Mesitylene Sulfonylamide, 3: Compound 102 (2.12 g, 20.78 mmol) was taken into 14 ml of dioxane/water (1:1) and a solution of mesitylenesulfonyl chloride (11.82 g, 54.05 mmol) in 7 ml dioxane was added dropwise (in about 0.5 h)

while maintaining the pH at about 12 with 5% NaOH solution. A thick gum-like product separated out of solution. Supernatant liquid was decanted. Crude product was column purified with hexanes and ethyl acetate (4:1) to afford 3.37 g (35%) of product 3 as white solid, m.p. 101–102° C. ¹H NMR (CDCl₃): δ 0.95 (t, J=6.99 Hz, 3H, CH₃), 1.68 (pent, J=6.25 Hz, 2H, NCCH₂), 2.29 (s, 6H, 2CH₃), 2.56 (s, 6H, 2CH₃), 2.62 (s, 6H, 2CH₃), 2.91 (m, 2H, NHCH₂), 3.12 (q, J=7.0 Hz, 2H, NCH₂), 3.31 (t, J=6.44 Hz, 2H, NCH₂), 4.93 (br, NH), 6.93 (s, 2H aromatic), 6.95 (s, 2H, aromatic); ¹³C NMR (CDCl₃): δ 12.57, 20.93, 22.81, 22.89, 27.84, 39.26, 39.99, 42.29, 131.95, 132.05, 133.32, 138.90, 140.00, 142.02, 142.57.

Compound 15' was prepared according to the method of Ashton et al. (1988), *J. Med. Chem.* 31, 2304.

Mesitylene sulfonic acid 2-mesitylene sulfonyloxymethyl-trans-cyclopropyl methyl ester, 16': Compound 15' (1.65 g, 16.18 mmol) was dissolved in 10 ml of pyridine and cooled to 0° C. A solution of mesitylenesulfonyl chloride (9.13 g, 41.75 mmol) in 23 ml of pyridine was added slowly over a span of about 15 minutes, stirred at room temperature for 3 hours, and then poured onto ice (75 g). White solid precipitated out which was filtered and recrystallized from EtOH; yield 2.8 g (37%); m.p. 83–84° C. ¹H NMR (CDCl₃): δ 0.50–0.65 (m, 2H, CyPrCH₂), 1.00–1.20 (m, 2H, CyPrCH), 2.31 (s, 3H, CH₃), 2.34 (s, 3H, CH₃), 2.60 (s, 6H, 2CH₃), 2.61 (s, 6H, 2CH₃), 3.70–3.90 (m, 4H, 2CH₂), 6.97 (s, 2H, aromatic), 6.99 (s, 2H, aromatic); ¹³C NMR (CDCl₃): δ 9.47, 16.36, 21.04, 22.57, 72.13, 130.62, 131.76, 139.79, 143.37. MS: (m/z) 466 (M⁺), 452, 386, 200, 171, 134, 118 (100%), 91, 77.

N-Ethyl-N-{3[(2-{[3-ethyl-mesitylenesulfonylamino]propyl}-mesitylenesulfonylamino)methyl]-(E)-cyclopropylmethyl)-mesitylenesulfonylamino]propyl}mesitylenesulfonylamide, 22': Compound 3 (2.5 g, 5.37 mmol) was taken into a flame dried 3-necked flask, dissolved in 40 ml anhydrous DMF. NaH (95%, 300 mg) was added slowly under argon atmosphere at 32° C. Contents were stirred at room temperature for 0.5 hours, then a solution of 16' (1.16 g, 2.49 mmol) in 35 ml anhydrous DMF was added over a period of 10 minutes. The reaction mixture was heated to 70° C., stirred for 4 hrs, cooled to 0° C., followed by quenching with 10 ml H₂O. The quenched reaction was then extracted with ether (3×30 ml), the combined organic layers were washed with H₂O (30 ml×4) and brine (2×25 ml). The solvents were then removed to afford crude oil, which was purified by column chromatography to yield 22', a low melting white semi-solid in 40% yield (1.0 g). ¹H NMR (CDCl₃): δ 0.34 (dd, J=6.6, 6.8 Hz, 1H, CyPrCH), 0.70–0.80 (m, 1H, CyPr), 0.95 (t, J=7.1 Hz, 6H, 2CH₃), 1.60–1.75 (m, 4H, 2CH₂), 2.30 (s, 12H, 4CH₃), 2.54 (s, 24H, 8CH₃), 2.60–2.89 (m, 2H, NCH), 2.93–3.25 (m, 14H, NCH₂), 6.92 (s, 4H, aromatic), 6.93 (s, 4H, aromatic); ¹³C NMR (CDCl₃): δ 12.69, 15.69, 20.95, 22.75, 25.29, 40.17, 42.64, 43.40, 48.90, 131.92, 131.98, 132.99, 133.26, 140.08, 140.12, 142.34, 142.49.

N-Ethyl-N'-(2-(3'-ethylamino-propylaminomethyl)-trans-cyclopropylmethyl)-propane-1,3-diamine tetrahydrochloride, 23: Phenol (1.68 g, 17.8 mmol) and HBr (33%) in glacial acetic acid (8.9 mL) were added successively to a solution of 22' (447 mg, 0.45 mmol) in CH₂Cl₂ (5 mL) at room temperature. The solution was stirred for 48 hrs. H₂O (6 mL) was added, followed by extraction with CH₂Cl₂ (3×8 mL). The aqueous layer was evaporated under reduced pressure and the residue was taken up in 10N NaOH (3 mL), followed by extraction with CHCl₃ (12×6 mL). Chloroform was distilled off to afford a thick gum which was

taken into anhydrous ether. Anhydrous HCl gas was passed through the solution to precipitate the tetrahydrochloride of 23 as white solid. (140 mg, 75% yield). The product was recrystallized from ethanol/ether; mp=270° C. (dec). ¹H NMR (D₂O): 80.88 (dd, J=6.8, 7.0 Hz, 2H, CyPrCH₂), 1.25 (t, J=6.1 Hz, 2H, CyPrCH), 1.30 (t, J=7.3 Hz, 6H, 2CH₃), 2.12 (pent., J=7.9 Hz, 4H, NCCH₂), 2.89 (dd, J=8.5, 12.1 Hz, 2H, NCH₂), 3.08–3.30 (m, 14H, NCH₂); ¹³C NMR (D₂O): 813.62, 14.10, 17.25, 26.35, 46.70, 47.49, 47.81, 54.41; MS (m/z) 271.4 (M⁺H⁺), 307.3 (M⁺H⁺+2 HCl), 156.7, 136.3 (100%).

Compound 24: Compound 24 was prepared by the method of Ashton et al. (1988) in 75% yield as a colorless oil. ¹H NMR (CDCl₃) 80.22 (dt, J=5, 5 Hz, 1H, CyPrCH₂), 0.70–0.85 (m, 1H, CyPrCH₂), 1.35 (m, 2H, CyPrCH), 3.27 (dd, J=11, 11 Hz, 2H, CH₂OH), 3.70–3.90 (br, 2H, OH), 4.14 (dd, J=6, 11 Hz, 2H, CH₂OH); ¹³C-NMR (CDCl₃) 88.39, 17.43, 63.80.

Compound 25: Compound 25 was obtained using the method of Fabiano et al. (1987). To a solution of the diol 24 (1.7 g, 17.01 mmol) in anhydrous THF (10 mL) was added a 0.4 M solution of HN₃ in toluene (102 mL) followed by a solution of diisopropyl azidodicarboxylate (7.48 g, 36.99 mmol) in THF (15 mL). To the resulting mixture was added a solution of PPh₃ (21.6 g, 92.35 mmol) in THF (30 mL) with stirring. As the reaction is exothermic, the temperature was controlled by the rate of addition of the PPh₃ solution. Stirring was continued for 1 hour at RT and then overnight at 50° C. H₂O (3.5 mL) was added, stirred at 50° C. for another 6 hours. The solvents were removed under vacuum and the residue was partitioned between CH₂Cl₂ (100 mL) and 1N HCl (100 mL). The aqueous phase was extracted with CH₂Cl₂ (4×100 mL). Removal of H₂O under reduced pressure (40° C.) gave 25 as a white solid which was recrystallized from methanol/ether (0.650 g, 25%). mp=230° C. (dec); ¹H NMR (D₂O) 80.40–0.53 (m, 1H, CyPrCH), 1.00–1.15 (m, 1H, CyPrCHH), 1.30–1.48 (m, 2H, CyPrCH), 2.75–2.95 (m, 2H, NCH₂), 3.20–3.35 (m, 2H, NCH₂).

Compound 26: Compound 26 was obtained (93%) from 25 following the procedure 7. The product was purified by column chromatography using silica gel and hexanes:ethylacetate (3:2) as an eluant. mp=152–153° C.; ¹H NMR (CDCl₃) 80.06–0.1 (m, 1H, CyPrCH₂), 0.62–0.75 (m, 1H, CyPrCHH), 1.00–1.16 (m, 2H, CyPrCH), 2.29 (s, 6H, 2CH₃), 2.62 (s, 12H, 4CH₃), 2.69 (dd, J=5.0 and 8.6 Hz, 2H, NCHH), 3.04 (dd, J=5.0 and 13.2 Hz, NCHH), 3.70 (s, 2H, NH), 6.04 (s, 4H, aromatic).

Compound 27: Compound 27 was obtained from 26 in 64% yield; mp=56–58° C.; ¹H NMR (CDCl₃) 80.05–0.09 (m, 1H, CyPrCHH), 0.7–0.81 (m, 1H, CyPrCHH), 0.95 (t, J=7.1 Hz, 8H, 2CH₃ and 2H, CyPrCH), 1.65–1.79 (m, 4H, NCHH₂), 2.28 (s, 6H, 2CH₃), 2.29 (s, 6H, 2CH₃), 2.54 (s, 12H, 4CH₃), 2.55 (s, 12H, 4CH₃), 2.69–2.90 (m, 2H, NCHH), 3.00–3.20 (m, 12H, NCR), 3.29–3.40 (m, 2H, NCHH), 6.92 (s, 4H, aromatic), 6.93 (s, 4H, aromatic); ¹³C-NMR (CDCl₃) 812.63, 13.92, 20.96, 22.73, 22.77, 25.50, 40.07, 42.65, 43.49, 45.17, 131.92, 133.00, 140.06, 142.35; MS (m/z) 8999.40 (M⁺), 815.30, 533.20, 167.10, 119.00 (100%).

Compound 28 (Scheme 5): Compound 28 was prepared from 27 in 75% yield; mp=240° C. (dec); ¹H NMR (D₂O) 80.38–0.48 (m, 1H, CyPrCHH), 0.9–1.1 (m, 1H, CyPrCHH), 1.29 (t, J=6.3 Hz, 8H, 2CH₃ and 2H, CyPrCH), 1.98–2.10 (m, 4H, NCCH₂), 2.88–3.05 (m, 8H, NCH₂), 3.05–3.18 (m, 8H, NCH₂); ¹³C-NMR (D₂O) 812.63, 13.53, 15.74, 26.89, 45.82, 47.30, 47.39, 50.58; HPLC R_T=26.97.

Compound 28 (via Scheme 5A): Referring specifically to Scheme 5A, compound 24 was made as described above. Mesitylenesulfonic acid 2-mesitylenesulfonyloxymethyl-cis-cyclopropylmethyl ester, 25' was obtained from 24 as described above for compound 16'. mp. 240° C. ¹H NMR (CDCl₃): 80.31–0.40 (m, 1H, CyPr), 0.86–0.99 (m, 1H, CyPr), 1.25–1.41 (m, 2H, CyPr), 2.32 (s, 6H, 2CH₃), 2.60 (s, 12H, 4CH₃), 3.85–4.10 (m, 4H, 2CH₂), 6.99 (s, 4H, aromatic); ¹³C NMR (CDCl₃): 89.71, 15.25, 21.08, 22.61, 69.61, 130.63, 131.79, 139.82, 143.39; MS 466 (M⁺), 452, 266, 200, 185, 171, 119 (100%).

N-Ethyl-N'-{3[(2-{[3-ethyl-mesitylenesulfonylamino]propyl}-mesitylenesulfonylamino)methyl]-(Z)-cyclopropylmethyl)-mesitylenesulfonylamino]-propyl}mesitylenesulfonylamide, 27, was obtained by reacting 25' with 3 in 64% yield; mp=56–58° C. ¹H NMR (CDCl₃): 80.05–0.09 (m, 1H, CyPrCHH), 0.7–0.81 (m, 1H, CyPrCHH), 0.95 (t, J=7.1 Hz, 8H, 2CH₃ and 2H, CyPrCH), 1.65–1.79 (m, 4H, NCHH₂), 2.28 (s, 6H, 2CH₃), 2.29 (s, 6H, 2CH₃), 2.54 (s, 12H, 4CH₃), 2.55 (s, 12H, 4CH₃), 2.69–2.90 (m, 2H, NCHH), 3.00–3.20 (m, 12H, NCH₂), 3.29–3.40 (m, 2H, NCHH), 6.92 (s, 4H, aromatic), 6.93 (s, 4H, aromatic); ¹³C NMR (CDCl₃): 812.63, 13.92, 20.96, 22.73, 22.77, 25.50, 40.07, 42.65, 43.49, 45.17, 131.92, 133.00, 1460.06, 142.35; MS (m/z) 8999.40 (M⁺), 815.30, 533.20, 167.10, 119.00 (100%).

N-Ethyl-N'-(2-(3'-ethylamino-propylaminomethyl)-cis-cyclopropylmethyl)-propane 1,3-diamine tetrahydrochloride, 28: This compound was synthesized by following the procedure described above for compound 23 (Scheme 4A) in 75% yield. White solid, mp=240° C. (dec). ¹H NMR (D₂O): 80.38–0.48 (m, 1H, CyPrCHH), 0.9–1.1 (m, 1H, CyPrCHH), 1.29 (t, J=6.3 Hz, 8H, 2CH₃ and 2H, CyPrCH), 1.98–2.10 (m, 4H, NCCH₂), 2.88–3.05 (m, 8H, NCH₂), 3.05–3.18 (m, 8H, NCH₂); ¹³C NMR (D₂O): 812.63, 13.53, 15.74, 26.89, 45.82, 47.30, 47.39, 50.58; HPLC R_T=26.97.

Compound 31: This compound was prepared by the method of Buchman et al. (1942). Mesitylenesulfonyl chloride (4.55 g, 21 mmol) in dioxane (15 mL) was added dropwise with magnetic stirring to 29 (1.1 g, 6.9 mmol) in 20 mL of a mixture of NaOH:dioxane (2:1). The mixture was maintained at pH 10–11 with occasional addition of a 5% solution of KOH. Once the addition was completed, the reaction mixture was left for an additional hour. The white solid was filtered off and dried to afford 31 (2 g, 65%). mp 201–202° C. (crystallized from methanol-water); ¹H NMR (CDCl₃) 81.35 (m, 2H), 1.95 (m, 2H), 2.30 (s, 6H), 2.60 (s, 6H), 2.60 (s, 12H), 3.32 (m, 2H), 5.18 (br, 2H), 6.95 (s, 4H); ¹³C-NMR (CDCl₃) 820.96, 22.88, 23.80, 54.68, 132.09, 134.10, 139.16, 142.80. MS (m/z, relative intensity), 451 (MH⁺, 2), 267 (73), 183 (48).

Compound 32: Compound 32 (70%) was obtained from 30 following the procedure described by Buchman et al. (1942). mp=192–193° C. (crystallized from methanol-water) ¹H NMR (CDCl₃) 81.90–2.18 (m, 4H), 2.35 (s, 6H), 2.60 (s, 12H), 3.63 (br, 2H), 5.41 (br, 2H), 6.96 (s, 4H); ¹³C-NMR (CDCl₃) δ: 20.95, 22.89, 25.32, 51.20, 132.04, 134.08, 139.30, 142.51 MS (m/z, relative intensity) 451 (MH⁺, 2), 267(76), 183(65).

Compound 33: NaH (95%, 115 mg, 4.8 mmol) was added to a solution of 31 (900 mg, 2 mmol) in DMF (20 mL) at 0° C. The mixture was stirred at 0° C. for 30 minutes and a solution of the alkylating agent 9 (1.53 g, 4.4 mmol) in DMF (10 mL) was added dropwise. The reaction mixture was stirred at 0° C. for 30 minutes and then at 70–80° C.

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overnight, cooled and quenched with water (5 mL). The solvent was distilled off with a vacuum pump and the residue was extracted into chloroform-water. The organic layer was dried and evaporated to leave a residue that was purified by flash chromatography on silica gel (elution with 20% ethyl acetate in hexanes) to yield 1.2 g (60%) of 33 as a colorless, glassy oil. ¹H-NMR (CDCl₃) δ: 0.98 (t, 6H), 1.40–1.70 (b, 6H), 1.82 (b, 2H), 2.28, 2.30 (s, 12H), 2.56–2.58 (s, 24H), 2.80–3.20 (m, 12H), 4.20 (br, 2H), 6.93–6.96 (s, 8H); ¹³C-NMR (CDCl₃) δ: 12.70, 20.94, 21.70, 22.76, 22.90, 28.20, 40.20, 41.54, 43.16, 55.48, 131.91, 132.10, 133.02, 133.18, 140.23, 142.80, 143.17, FABMS (m/z, relative intensity) 985.4 (MH⁺, 37), 801.4 (84), 619.3 (20), 619.3 (21).

Compound 34: Compound 34 (50%) was obtained from 32 following the procedure described above. ¹H-NMR (CDCl₃) δ: 0.98 (t, 6H), 1.50 (m, 4H), 1.80 (br, 2H), 2.12 (br, 2H), 2.30 (s, 6H), 2.52, 2.54 (s, 12H), 2.82 (m, 4H), 2.93–3.23 (m, 8H), 4.20 (br, 2H), 6.93 (s, 8H); ¹³C-NMR (CDCl₃) δ: 12.69, 20.94, 22.75, 22.82, 24.63, 25.94, 40.14, 42.16, 43.25, 55.43, 131.89, 132.23, 133.51, 135.08, 139.52, 140.16, 142.22, 142.88; FABMS (m/z, relative intensity) 985.4 (MH⁺, 11%), 801.3 (59), 619.3 (26).

Compound 35: Phenol (800 mg, 8.5 mmol) and 30% HBr in glacial acetic acid (14 mL) were added successively to a solution of the tetramide 33 (300 mg, 0.30 mmol) in CH₂Cl₂ (14 mL) at room temperature. The solution was stirred at room temperature for 48 hrs. H₂O (20 mL) was added followed by extraction with CHCl₃ (3×20 mL). The aqueous portion was evaporated to dryness and the residue was taken up in 10N NaOH (5 mL) followed by extraction with CHCl₃ (10×5 mL). The combined organic phases were dried and evaporated. The resulting residue was dissolved in ethanol (3 mL) and acidified with three drops of concentrated HCl, cooled and filtered. White crystals of 35 were obtained (50 mg, 40%). ¹H-NMR (D₂O) δ: 1.20 (t, 6H), 1.80–2.21 (m, 6H), 2.35 (m, 2H), 3.08 (m, 12H), 4.00 (m, 2H); ¹³C-NMR (D₂O) δ: 14.20, 22.57, 26.34, 46.26, 46.76, 47.44, 57.57, MS (m/z, relative intensity) 257.2 (MH⁺, 27), 293.3 (M+2H⁺+2Cl⁻, 4), 172.2 (100); HPLC: R_f=27.47 min.

Compound 36: Compound 36 (40 mg, 37%) was obtained following the procedure described above for 35 using 300 mg of 34 as a reactant. ¹H-NMR (D₂O) δ: 2.10–2.25 (m, 4H), 2.35–2.55 (m, 4H), 3.08–3.30 (m, 12H), 4.15 (m, 2H); ¹³C-NMR (D₂O) δ: 14.12, 25.04, 26.28, 46.72, 47.31, 47.46, 57.98; ESIMS (m/z, relative intensity): 257.3 (MH⁺, 100), 293.3 (M⁺ 2H⁺+2Cl⁻, 6), 172.3 (12); HPLC: R_f=25.91 min.

Compound 39: Compound 39 was obtained from 37 (see Buchman et al. (1942)) following the procedure described above. From 3.85 g (22 mmol) of 37 was obtained 1.68 g (65%) of 39 as a colorless oil. ¹H-NMR (CDCl₃) δ: 1.60 (m, 2H), 1.85 (m, 2H), 2.20 (br, 2H), 3.38 (m, 2H), 3.60 (m, 2H); ¹³C-NMR (CDCl₃) δ: 20.80, 43.46, 66.33.

Compound 40: A three-necked flask equipped with an addition funnel and a condenser containing LiAlH₄ (2.8 g, 73.6 mmol) was cooled in an ice bath as 80 mL of dry THF was added under a nitrogen atmosphere. The ice bath was removed and the mixture was stirred at room temperature while a solution of 38 (see Buchman et al. (1942)) (6.16 g 36 mmol) in 30 mL of THF was added dropwise. The mixture was stirred at reflux for 18 hrs., After cooling in an ice bath, the mixture was treated cautiously with a saturated solution of NH₄Cl (16 mL) and then with ethyl acetate (16 mL). The insoluble salts were removed by filtration and the filtrate was evaporated under reduced pressure. The residue was dissolved in ethyl acetate, dried (NaSO₄), filtered and

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evaporated to give 3.6 g (86%) of a colorless oil. ¹H-NMR (CDCl₃) δ: 1.58 (m, 2H), 2.05 (m, 2H), 2.70 (br, 2H), 3.60 (m, 2H), 3.82 (m, 2H); ¹³C-NMR (CDCl₃) δ: 20.32, 20.82, 37.46, 62.72.

Compound 41: Compound 41 was obtained from 39 following the procedure described immediately above. From 1.97 g (17 mmol) of 39 was obtained 700 mg (21%) of crude 41 which was used in subsequent steps without further purification.

Compound 42: To a solution of the alcohol 40 (1.97 g, 17 mmol) in THF (10 mL) was added 42 mL of a 0.95 M solution of hydrazoic acid (HN₃, 39.8 mmol), followed by a solution of diisopropyl-azodicarboxylate (7.45 g, 69 mmol) in THF (20 mL). To the resulting mixture, a solution of triphenylphosphine (19.6 g, 75 mmol) in THF (85 mL) was added while stirring. After 1 hour at room temperature the reaction mixture was heated at 50° C. for 6 hours, the solvents were removed in vacuo and the residue was partitioned between CH₂Cl₂ (60 mL) and 1 N HCl (30 mL). The aqueous phase was extracted with additional CH₂Cl₂ (3×10 mL). Removal of the water under reduced pressure gave the diamino hydrochloride 42 (800 mg, 25%). This was used in subsequent steps without further purification.

Compound 44: Mesitylenesulfonylchloride (2.70 g, 126 mmol) in dioxane (10 mmol) was added dropwise with stirring to 42 (800 mg, 4.2 mmol) dissolved in a mixture of 1N NaOH and dioxane (1:1). The mixture was maintained at pH 10–11 with occasional addition of a 5% solution of iN NaOH. Once the addition was complete, the reaction mixture was stirred for an additional hour. The solvent was decanted and the oily residue was washed with hexane, filtered, and dried, which afforded 490 mg (25%) of 44. mp=138–139° C. (crystallized from methanol-water). ¹H-NMR (CDCl₃) δ: 1.50 (m, 2H), 1.88 (m, 2H), 2.30 (br, 8H), 2.62 (s, 12H), 2.92 (m, 4H), 5.05 (t, 2H), 6.90 (s, 4H); ¹³C-NMR (CDCl₃) δ: 20.92, 21.65, 22.95, 39.01, 46.82, 131.96, 134.20, 139.03, 142.12; MS (m/z, relative intensity): 478 (M⁺, 12), 295 (90).

Compound 43: Compound 43 was obtained from 41 following the procedure described above. From 700 mg of 41 was obtained 400 mg (25%) of 43. mp=130–132° C. (crystallized from methanol-water). ¹H-NMR (CDCl₃) δ: 1.50 (m, 2H), 1.83 (m, 2H), 2.25 (m, 2H), 2.30 (s, 6H), 2.60 (s, 12H), 2.90 (m, 4H), 5.20 (t, 2H), 6.95 (s, 4H); ¹³C-NMR (CDCl₃) δ: 20.91, 21.66, 22.95, 38.97, 46.80, 131.95, 133.72, 139.03, 142.11; MS (m/z, relative intensity): 478 (M⁺, 7), 295 (54).

Compound 46—NaH (95%, 52.3 mg, 2.18 mmol) was added to a solution of 44 (400 mg, 0.84 mmol), in DMF (20 mL) at 0° C. The mixture was stirred at 0° C. for 30 minutes, and a solution of N-ethyl-N-mesitylenesulfonamide-3-bromopropylamine 9 (758 mg, 218 mmol) in DMF (15 mL) was added dropwise. The reaction mixture was then stirred at 0° C. for 15 minutes, and then at 70–80° C. overnight, and then cooled and cautiously quenched with water (25 mL). The mixture was extracted with ethyl ether (3×30 mL), and the pooled organic fractions were washed with H₂O (4×30 mL) and brine (2×20 mL), and dried (NaSO₄). The solvent was evaporated under reduced pressure, yielding an oil. The crude oil was purified by column chromatography using hexanes:ethyl acetate (8:2), followed by an 8:3 hexanes:ethyl acetate eluant. 800 mg (94%) of 46 were obtained as a glassy oil. ¹H-NMR (CDCl₃) δ: 0.95 (t, 6H), 1.30 (m, 3H), 1.61 (m, 5H), 2.10 (m, 2H), 2.32 (s, 12H), 2.52 (s, 24H), 2.80–3.35 (m, 14H), 3.30 (m, 2H), 6.95, 6.98 (s, s, 8H); ¹³C-NMR (CDCl₃) δ: 12.64,

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20.93, 22.72, 22.82, 23.30, 24.87, 37.46, 39.97, 42.47, 43.60, 50.06, 131.91, 133.70, 139.99, 142.30; FABMS (*m/z*, relative intensity): 1013.4 (*M*⁺, 33), 829.5 (29).

Compound 45: Compound 45 was obtained from 43 following the procedure described immediately above. From 300 mg of 43 was obtained 500 mg (78%) of a glassy oil product. ¹³C-NMR (CDCl₃) δ: 12.63, 20.93, 22.72, 22.81, 23.28, 24.97, 37.45, 39.95, 42.29, 43.68, 50.01, 131.91, 133.70, 139.97, 140.03, 142.31; FABMS (*m/z*, relative intensity): 1013.5 (*M*⁺, 28) 829.5 (28).

Compound 48 (Scheme 9): Phenol (3.17 g, 33.7 mmol) and 30% HBr in glacial acetic acid (17 mL) were added successively to a solution of 46 (759 mg 0.74 mmol) in CH₂Cl₂ (10 mL) at room temperature. The solution was stirred for 48 hours, H₂O (10 mL) was added, followed by extraction with CH₂Cl₂ (3×10 mL). The aqueous layer was evaporated under reduced pressure and the residue was taken up in 10N NaOH (5 mL) followed by extraction with CHCl₃ (12×10 mL). After removal of CHCl₃, the residue was taken up in ethanol (10 mL) and acidified with concentrated HCl (0.5 mL). The precipitate was recrystallized from ethanol-ether to give 190 mg (60%) of 48 as white crystals. ¹H NMR (D₂O), δ: 1.30 (t, 6H), 1.85 (m, 2H), 2.08–2.25 (m, 6H), 2.50 (br, 2H), 3.10–3.30 (m, 16H); ¹³C-NMR (D₂O) δ: 14.24, 26.44, 38.77, 46.80, 47.60, 48.41, 55.05; HPLC: *R*_T=28.06 min.

Compound 47 (Scheme 8): Following the procedure described for 48, 50 mg (23%) of 47 were obtained as white crystals. ¹H NMR (D₂O) δ: 1.30 (t, 6H), 1.80 (m, 2H), 2.02–2.28 (m, 6H), 2.50 (br, 2H), 3.02–3.30 (m, 16H); ¹³C-NMR (D₂O) δ: 14.22, 26.43, 38.77, 46.82, 47.60, 48.40, 55.03; HPLC: *R*_T=28.047 min.

Compound 47 (via Scheme 8A): Referring specifically to Scheme 8A, mesitylenesulfonic acid 2-mesitylenesulfonyloxymethyl-trans-cyclobutyl methyl ester, 41' (83%) was obtained from 39 following the procedure described for 16' (Scheme 4A). m.p. 77–78° C. ¹H NMR (CDCl₃): δ: 1.65 (m, 2H), 1.95 (m, 2H), 2.30 (s, 6H), 2.58, 2.60 (s, s 12H), 3.88 (d, 4H), 6.96 (s, 4H); ¹³C NMR (CDCl₃): δ: 20.67, 20.86, 22.39, 36.37, 71.28, 130.35, 131.59, 131.62 m 139.60, 143.17. EI-MS *m/z* (relative intensity) 48 (*M*⁺, 8), 281 (5), 199 (22), 183 (36).

N-Ethyl-N'-{3[(2-{[3-ethyl-mesitylenesulfonylamino]propyl]-mesitylenesulfonylamino-methyl)-trans-cyclobutylmethyl]-mesitylenesulfonylamino]-propyl]mesitylenesulfonylamide, 45, was obtained by reacting 41' with 3 following the procedure described for the synthesis of 22' in 78% yield as a glassy oil. ¹H NMR (CDCl₃): δ: 13C NMR (CDCl₃): δ: 12.63, 20.93, 22.72, 22.81, 23.28, 24.97, 37.45, 39.95, 42.29, 43.68, 50.01, 131.91, 133.70, 139.97, 140.03, 142.31; FABMS (*m/z*, relative intensity): 1013.5 (*M*⁺, 28) 829.5 (28).

N-Ethyl-N'-(2-(3'-ethylamino-propylaminomethyl)-trans-cyclobutylmethyl)-propane-1,3-diamine tetrahydrochloride, 47: Following the procedure described above for 23 (Scheme 4A), 50 mg (23%) of 47 were obtained as white solid. ¹H NMR (D₂O): δ: 1.30 (t, 6H), 1.80 (m, 2H), 2.02–2.28 (m, 6H), 2.50 (br, 2H), 3.02–3.30 (m, 16H); ¹³C NMR (D₂O): δ: 14.22, 26.43, 38.77, 46.82, 47.60, 48.40, 55.03; HPLC: *R*_T=28.047 min.

Compound 48 (Scheme 9A): referring specifically to Scheme 9A, mesitylenesulfonic acid 2-mesitylene sulfonyloxymethyl-cis-cyclobutylmethyl ester, 42' (80%) was obtained from 40 following the procedure described for 16' m.p. 92–93° C. ¹H NMR (CDCl₃) δ: 1.72 (m, 2H), 2.05 (b, 2H), 2.31 (s, 6H), 2.57, 2.59 (s, 12H), 2.78 (b, 2H),

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3.85–4.10 (m, 4H), 6.96 (s, 4H). ¹³C NMR (CDCl₃) δ: 20.89, 21.49, 22.41, 34.94, 69.14, 130.41, 131.66, 139.62, 139.65, 143.20, 143.25. EI-MS *m/z* (relative intensity) 480 (*M*⁺, 0.7), 281 (7), 199 (22), 183 (47).

N-Ethyl-N'-(2-(3'-ethylamino-propylaminomethyl)-cis-cyclobutylmethyl)-propane-1,3-diamine tetrahydrochloride, 48: This compound was synthesized according to the procedure described for 23 (Scheme 4A) in 60% yield as a white solid. The product was recrystallized from ethanol-ether to afford 190 mg (60%) of 48 as white crystals. ¹H NMR (D₂O): δ: 1.30 (t, 6H), 1.85 (m, 2H), 2.08–2.25 (m, 6H), 2.50 (b, 2H), 3.10–3.30 (m, 16H); ¹³C NMR (D₂O): δ: 14.24, 26.44, 38.77, 46.80, 47.60, 48.41, 55.05; HPLC: *R*_T=28.06 min.

Compound 50: This compound was prepared according to the method of Miller et al. (1959). Using 49 as a reactant, 50 was prepared in 70% yield. ¹H NMR (CDCl₃) δ: 1.10 (s, slightly broad, 4H, CH₂OH), 5.83 (s, again slightly broad, 2H, CH=CH); ¹³C-NMR (CDCl₃, MeOH) δ: 62.07, 130.08.

Compound 51: Compound 51 (86%) was prepared as described above for 25 as a white solid. ¹H NMR (D₂O) δ: 3.69 (d, J=5 Hz, 4H, NCH₂), 6.00–6.10 (m, 2H, CH=CH); ¹³C-NMR (D₂O) δ: 43.10, 130.75.

Compound 52: Compound 52 (45%) was prepared as described above for 25 from commercially available cis-butene-1,4-diol 50'. mp=205° C. (dec); ¹H NMR (D₂O) δ: 3.77 (d, J=5.5 Hz, 4H, NCH₂), (t, J=4.6 Hz, 2H, CH=CH), ¹³C-NMR (D₂O) δ: 38.90, 129.74.

Compound 53: Compound 53 was prepared from 51 in 29% yield mp=171–172° C.; ¹H NMR (CDCl₃) δ: 3.31 (s, 6H, 2CH₃), 2.60 (s, 12H, 4CH₃), 3.45–3.55 (m, 4H, NCH₂), 5.40–5.50 (m, 2H, CH=CH), 6.95 (s, 4H, aromatic).

Compound 54: Compound 54 was prepared from 52 in 74% yield. mp=109–110° C.; ¹H NMR (CDCl₃) δ: 3.31 (s, 6H, 2CH₃), 2.60 (s, 12H, 4CH₃), 3.45–3.55 (m, 4H, NCH₂), 5.40–5.50 (m, 2H, CH=CH), 6.95 (s, 4H, aromatic); ¹³C-NMR (CDCl₃) δ: 20.95, 22.92, 39.05, 128.30, 132.03, 133.62, 139.11, 142.42; MS (*m/z*) 451 (*M*⁺), 267, 251, 183, 119 (100%), 91; Anal: (C₂₂H₃₀N₂O₄S₂).

Compound 55: Compound 55 was prepared in 91% yield as described above for 10. mp=135–136° C.; ¹H NMR (CDCl₃) δ: 80.95 (t, J=7.1 Hz, 6H, 2CH₃), 1.59–1.70 (m, 4H, NCH₂), 2.29 (s, 12H, 4CH₃), 2.53 (s, 12H, 4CH₃), 2.55 (s, 12H, 4CH₃), 2.99 (t, J=7.3 Hz, 8H, NCH₂), 3.08 (q, J=7.1 Hz, 4H, NCH₂), 3.65 (d, J=5 Hz, 4H, C=C–CH₂), 5.40–5.50 (m, 2H, CH=CH), 6.92 (s, 4H, aromatic), 6.93 (s, 4H, aromatic); ¹³C-NMR (CDCl₃) δ: 12.62, 20.90, 22.68, 22.77, 25.56, 40.02, 42.53, 43.79, 43.83, 128.51, 131.88, 132.00, 132.71, 133.00, 140.02, 140.17, 142.30, 142.66.

Compound 56: Compound 56 was prepared as described above for 10 from 54 in 88% yield. mp=74–75° C.; ¹H NMR (CDCl₃) δ: 80.93 (t, J=7.1 Hz, 6H, 2CH₃), 1.60–1.70 (m, 4H, NCH₂), 2.29 (s, 6H, 2CH₃), 2.30 (s, 6H, 2CH₃), 2.52 (s, 12H, 4CH₃), 2.55 (s, 12H, 4CH₃), 2.95–3.02 (m, 8H, NCH₂), 3.07 (q, J=7.1 Hz, 4H, NCH₂), 3.74 (d, J=4.6 Hz, 4H, NCH₂), 6.90 (s, 4H, aromatic), 6.93 (s, 4H, aromatic); ¹³C-NMR (CDCl₃) δ: 12.69, 20.93, 22.71, 22.80, 25.20, 40.12, 42.56, 43.25, 47.06, 129.50, 131.93, 132.03, 132.86, 133.25, 140.07, 140.12, 142.36, 142.62; MS (*m/z*) 985.3 (*M*⁺), 801.3, 718.2, 617.2, 519.2, 336.1, 240.1, 183.0, 119 (100%).

Compound 57 (Scheme 10): Compound 57 was prepared as described above for 12 starting from 35 in 86% yield. mp=250° C. (dec); ¹H NMR (D₂O) δ: 1.30 (t, J=7.4 Hz, 6H, 2CH₃), 2.05–2.20 (m, 4H, NCH₂), 3.05–3.20 (m, 12H, NCH₂), 3.75–3.80 (m, 4H, NCH), 6.04–6.10 (m, 2H,

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CH=CH); ^{13}C -NMR (D_2O) δ 13.34, 25.57, 45.90, 46.71, 46.80, 51.04, 131.25; HPLC: R_T =27.03 min.

Compound 58 (Scheme 11): Compound 58 was prepared from 56 in 86% yield. mp=240° C.; ^1H NMR (D_2O) δ 1.30 (t, J=7.3 Hz, 6H, CH_3), 2.10–2.5 (m, 4H, NCCH_2), 3.05–3.25 (m, 12H, NCH_2), 3.87 (d, J=4.8 Hz, 4H, NCH_2), 5.98 (t, J=4.8 Hz, 2A, $\text{CH}=\text{CH}$); ^{13}C -NMR (D_2O) δ 13.35, 25.69, 45.93, 46.70, 46.96, 47.02, 129.31; HPLC: R_T =26.89 min.

Compound 57 (via Scheme 10A): Referring specifically to Scheme 10A, mesitylenesulfonic acid-4-mesitylenesulfonyloxy-E-but-2-enyl ester, 51': Diol 50 (1.76 g, 20 mmol), and benzytriethylammonium bromide (270 mg, 1 mmol) were taken in a mixture of 30 mL 50% KOH and 30 mL of dioxane. The reaction mixture was stirred in an ice water bath as mesitylene sulfonyl chloride (8.72 g, 40 mmol) in 30 mL of dioxane as added dropwise. When the addition was over, the stirring was continued for an additional hour. Excess water was added and after cooling a white precipitate was filtered. Crystallization from chloroform-hexane yielded 7.0 g 51' (77%); m.p. 119–120° C. ^1H NMR (CDCl_3): δ 2.35 (s, 6H), 2.60 (s, 12H), 4.45 (d 4H), 5.75 (b, 2H), 6.95 (s, 4H); ^{13}C NMR (CDCl_3): δ 20.96, 22.52, 67.96, 127.67, 131.69, 131.74, 139.79, 143.45. EI-MS m/z (relative intensity) 452 (M^+ , 1), 253 (17), 200 (81), 183 (39).

N-Ethyl-N-{3[(2-{[3-ethyl-mesitylenesulfonylamino]propyl}-mesitylenesulfonylamino)methyl]-but-(E)-2-enyl}-mesitylenesulfonylamino]-propyl}-mesitylenesulfonylamide 55: This compound was prepared in 91% yield as described above for 22'. mp=135–136° C. ^1H NMR (CDCl_3): δ 0.95 (t, J=7.1 Hz, 6H, 2CH_3), 1.59–1.70 (m, 4H, NCCH_2), 2.29 (s, 12H, 4CH_3), 2.53 (s, 12H, 4CH_3), 2.55 (s, 12H, 4CH_3), 2.99 (t, J=7.3 Hz, 8H, NCH_2), 3.08 (q, J=7.1 Hz, 4H, NCH_2), 3.65 (d, J=5 Hz, 4H, $\text{C}=\text{C}-\text{CH}_2$), 5.40–5.50 (m, 2H, $\text{CH}=\text{CH}$), 6.92 (s, 4H, aromatic), 6.93 (s, 4H, aromatic); ^{13}C NMR (CDCl_3): δ 12.62, 20.90, 22.68, 22.77, 25.56, 40.02, 42.53, 43.79, 43.83, 128.51, 131.88, 132.00, 132.71, 133.00, 140.02, 140.17, 142.30, 142.66.

N,N¹-bis(3-ethylaminopropyl)-(E)-but-2-ene-1,4-diamine tetrahydrochloride, 57, was prepared as described above for 23 (Scheme 4A) in 86% yield. mp=250° C. (dec). ^1H NMR (D_2O): δ 1.30 (t, J=7.4 Hz, 6H, 2CH_3), 2.05–2.20 (m, 4H, NCCH_2), 3.05–3.20 (m, 12H, NCH_2), 3.75–3.80 (m, 4H, NCH_2), 6.04–6.10 (m, 2H, $\text{CH}=\text{CH}$). ^{13}C NMR (D_2O): δ 13.34, 25.57, 45.90, 46.71, 46.80, 51.04, 131.25; HPLC: R_T =27.03 min.

Compound 58 (via Scheme 11A): Referring specifically to Scheme 11A, mesitylenesulfonic acid-4-mesitylenesulfonyloxy-E-but-2-enyl ester, 52' was obtained from 50' in 75% yield by following the procedure described above; m.p. 71–72° C. ^1H NMR (CDCl_3): δ 2.25 (s, 6H), 2.50 (s, 19H), 4.40 (d, 4H), 5.62 (b, 2H), 6.90 (s, 4H); ^{13}C NMR (CDCl_3): δ 20.90, 22.40, 63.66, 127.54, 131.71, 139.69, 143.46. EI-MS m/z (relative intensity) 452 (M^+ , 3), 253 (47), 199 (35), 183 (76).

N-Ethyl-N-{3[(2-{[3-ethyl-mesitylenesulfonylamino]propyl}-mesitylenesulfonylamino)methyl]-but-(Z)-2-enyl}-mesitylenesulfonylamino]-propyl}-mesitylenesulfonylamide, 56, prepared as described above for 22 (Scheme 4A) in 88% yield. mp=74–75° C. ^1H NMR (CDCl_3): δ 0.93 (t, J=7.1 Hz, 6H, 2CH_3), 1.60–1.70 (m, 4H, NCCH_2), 2.29 (s, 6H, 2CH_3), 2.30 (s, 6H, 2CH_3), 2.52 (s, 12H, 4CH_3), 2.55 (s, 12H, 4CH_3), 2.95–3.02 (m, 8H, NCH_2), 3.07 (q, J=7.1 Hz, 4H, NCH_2), 3.74 (d, J=4.6 Hz, 4H, NCH_2), 6.90 (s, 4H, aromatic), 6.93 (s, 4H, aromatic); ^{13}C NMR (CDCl_3): δ 12.69, 20.93, 22.71, 22.80, 25.20,

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40.12, 42.56, 43.25, 47.06, 129.50, 131.93, 132.03, 132.86, 133.25, 140.07, 140.12, 142.36, 142.62; MS (m/z) 985.3 (M^+).

N,N¹-bis(3-ethylaminopropyl)-(Z)-but-2-ene-1,4-diamine tetrahydrochloride, 58, was prepared from 56 in the same fashion as 23 (Scheme 4A) in 86.5% yield. mp=240° C. ^1H NMR (D_2O): δ 1.30 (t, J=7.3 Hz, 6H, CH_3), 2.10–2.59 (m, 4H, NCCH_2), 3.05–3.25 (m, 12H, NCH_2), 3.87 (d, J=4.8 Hz, 4H, NCH_2), 5.98 (t, J=4.8 Hz, 2H, $\text{CH}=\text{CH}$); ^{13}C NMR (D_2O): δ 13.35, 25.69, 45.93, 46.70, 46.96, 47.02, 129.31; HPLC: R_T =26.89 min.

Biological Activity Examples

These Examples are provided to illustrate the utility of the present compounds to inhibit neoplastic cell growth. As noted above, the Examples do not limit the scope of the invention described and claimed herein in any fashion.

Cell Lines and Media

Human breast cancer cell line MCF7 was grown in Richter's Improved Modified Eagle's Medium supplemented with 10% fetal bovine serum (FBS) and 2.2 g/L sodium bicarbonate. Human brain cancer cell line U251MG-NCI was grown in Dulbecco's Modified Eagle's Medium supplemented with 10% FBS. Human lung cancer cell line A549 was grown in Ham's F-12K medium (Fisher Scientific, Itasca, Ill.), supplemented with 10% FBS and 2 mM L-glutamine. Human colon cancer cell line HT29 was cultured in McCoy's 5A medium (Gibco, BRL, Gaithersburg, Md.), supplemented with 10% FBS. Human prostate cancer cell line PC3 was grown in Dulbecco's Modified Eagles Medium supplemented with 5% FBS. The A549 and MCF7 cell lines were cultured in 100 units/mL penicillin and 100 $\mu\text{g}/\text{mL}$ streptomycin. HT29 and U251MG cell lines were grown in 50 $\mu\text{g}/\text{mL}$ gentamycin. PC3 cell lines were maintained in 1% antibiotic-antimycotic solution (Sigma, St. Louis, Mo.). The cell cultures were maintained at 37° C. in 5% CO_2 /95% humidified air. All cell cultures are available from the American Type Culture Collection, Rockville, Md.

Examples 1–8

A standardized protocol was used to evaluate these test cultures and to generate the data shown in FIGS. 1–8:

Day 1:

Ten standard culture flasks for each drug to be tested were plated with 5×10^5 cells of a given type in 5 mL of media and allowed to incubate for 16–24 hours at 37° C.

Day 2:

Fresh stocks of the compounds to be evaluated are prepared. For each drug, two of the ten culture flasks prepared on Day 1 are used as controls. The control flasks are treated with solvent only. Four flasks for each compound are then treated with serially-diluted concentrations of the compound. The remaining flasks are left untouched. The cells are incubated for 4 hours at 37° C.

After 4' hours the control flasks are counted (2 counts for each flask) and the cells per mL calculated based on the average of the control counts. The cells are then re-plated into six 60 mm dishes for each flask from dilutions based on the cells/mL of the control. (In the various test runs, cell concentrations ranged from approximately 50 to approximately 800 cells per mL.)

Day 15–20:

The cells are monitored for colony formation. When visible, the cells are stained with 0.5% crystal violet (in 95% EtOH) and counted. The plating efficiency for each dish is

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then calculated. The plating efficiencies of the six dishes for each flask are averaged and the standard deviation is calculated. The fraction of cell survival at each concentration is determined based on the controls.

Example 1

In vitro effect of SL-11048 (Compound 57) on MCF7

Following the standard protocol described above, the effect of SL-11048 (Compound 57) on MCF7 cell lines was evaluated. The results are depicted in FIG. 1. As shown in FIG. 1, $ED_{50}=1.49 \mu M$.

Example 2

In vitro effect of SL-11038 (Compound 23) on MCF7

Following the standard protocol described above, the effect of SL-11038 (Compound 23) on MCF7 cell lines was evaluated. The results are depicted in FIG. 2. As shown in FIG. 2, $ED_{50}=1.34 \mu M$.

Example 3

In vitro effect of SL-11037 (Compound 28) on MCF7

Following the standard protocol described above, the effect of SL-11037 (Compound 28) on MCF7 cell lines was evaluated. The results are depicted in FIG. 3. As shown in FIG. 3, $ED_{50}=1.64 \mu M$.

Example 4

In vitro effect of SL-11043 (Compound 48) on MCF7

Following the standard protocol described above, the effect of SL-11043 (Compound 48) on MCF7 cell lines was evaluated. The results are depicted in FIG. 4. As shown in FIG. 4, $ED_{50}=1.64 \mu M$.

Example 5

In vitro effect of SL-11047 (Compound 58) on MCF7

Following the standard protocol described above, the effect of SL-11047 (Compound 58) on MCF7 cell lines was evaluated. The results are depicted in FIG. 5. As shown in FIG. 5, $ED_{50}=1.49 \mu M$.

Example 6

In vitro effect of SL-11044 (Compound 47) on MCF7

Following the standard protocol described above, the effect of SL-11044 (Compound 47) on MCF7 cell lines was evaluated. The results are depicted in FIG. 6. As shown in FIG. 6, $ED_{50}=1.79 \mu M$.

Example 7

In vitro effect of 10 μM concentrations of SL-11033 (13, ■), SL-11027 (12, ▲), SL-11034 (36, ▼), and SL-11028 (35, ◆) on U251MG-NCI cells.

Here, the above-identified compounds were administered in a 10 μM dose to cultures of the human brain cancer cell line U251MG-NCI and evaluated according to the standard protocol described above. The results are shown in FIG. 7. At the 10 μM dosage used, SL-11027 (12) displayed marked inhibition of cell growth. Control=●.

Example 8

In vitro effect of 40 μM concentrations of SL-11033 (13, ■), SL-11027 (12, ▲), SL-11034 (36, ▼), and SL-11028 (35, ◆) on U251MG-NCI cells.

This Example is identical to Example 7, with the exception that a 40 μM dose was administered. The results are depicted in FIG. 8. Here, at the 40 μM dosage used, SL-11034 (36) displayed marked inhibition of cell growth. Control=●.

Examples 9-20

Here, a conventional MTT assay was used to evaluate percent cell survival. Exponentially growing monolayer

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cells were plated in 96-well plates at a density of 500 cells per well and allowed to grow for 24 hours. Serial dilutions of the drugs were added to the wells. Six days after drug treatment, 25 μl of MTT solution (5 mg/ml) was added to each well and incubated for 4 hours at 37° C. Then 100 μl of lysis buffer (20% sodium dodecyl sulfate, 50% DMF, and 0.8% acetic acid, pH 4.7) was added to each well and incubated for an additional 22 hours. A microplate reader ("EMAX"-brand, Molecular Devices, Sunnyvale, Calif.) set at 570 nm was used to determine the optical density of the cultures. Results are expressed as a ratio of the optical density in drug-treated wells to the optical density in wells treated with vehicle only.

The ID_{50} doses for the compounds tested against the various cell lines are presented in Table 2. The ID_{50} is the drug concentration that killed 50% of the cultured cells.

TABLE 2

	Cytotoxic Activity Human Tumor Cell Lines				
	ID_{50} (μM)				
	A549	HT-29	PC-3	MCF7	U251MG NCI
SL-11037 (Cmpd 28)	0.12	1.6	7.4	>31.25	0.1
SL-11038 (Cmpd 23)	0.25	1.4	12.4	>31.25	0.1
SL-11043 (Cmpd 48)	0.1	1.5	>31.25	25.5	0.1
SL-11044 (Cmpd 47)	0.3	1.6	>31.25	>31.25	0.12
SL-11047 (Cmpd 58)	0.25	1.6	3.6	9.5	0.55
SL-11048 (Cmpd 57)	0.26	1.4	1.4	>31.25	2

Example 9

Using the standard MTT protocol described above, cultured HT29 cells were exposed to serial dilutions of compounds 28 (SL11037) and 23 (SL11038). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 9A and FIG. 9B. The ID_{50} for these compounds against HT29 is given in Table 2.

Example 10

Using the standard MTT protocol described above, cultured HT29 cells were exposed to serial dilutions of compounds 48 (SL11043) and 47 (SL11044). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 10A and FIG. 10B. The ID_{50} for these compounds against HT29 is given in Table 2.

Example 11

Using the standard MTT protocol described above, cultured HT29 cells were exposed to serial dilutions of compounds 58 (SL11047) and 57 (SL11048). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 11A and FIG. 11B. The ID_{50} for these compounds against HT29 is given in Table 2.

Example 12

Using the standard MTT protocol described above, cultured U251 MG cells were exposed to serial dilutions of

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compounds 28 (SL11037) and 23 (SL11038). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 12A and FIG. 12B. The ID₅₀ for these compounds against U251 MG is given in Table 2.

Example 13

Using the standard MTT protocol described above, cultured U251 MG cells were exposed to serial dilutions of compounds 48 (SL11043) and 47 (SL11044). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 13A and FIG. 13B. The ID₅₀ for these compounds against U251 MG is given in Table 2.

Example 14

Using the standard MTT protocol described above, cultured U251 MG cells were exposed to serial dilutions of compounds 58 (SL11047) and 57 (SL11048). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 14A and FIG. 14B. The ID₅₀ for these compounds against U251 MG is given in Table 2.

Example 15

Using the standard MTT protocol described above, cultured A549 cells were exposed to serial dilutions of compounds 28 (SL11037) and 23 (SL11038). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 15A and FIG. 15B. The ID₅₀ for these compounds against A549 is given in Table 2.

Example 16

Using the standard MTT protocol described above, cultured A549 cells were exposed to serial dilutions of compounds 48 (SL11043) and 47 (SL11044). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 16A and FIG. 16B. The ID₅₀ for these compounds against A549 is given in Table 2.

Example 17

Using the standard MTT protocol described above, cultured A549 cells were exposed to serial dilutions of compounds 58 (SL11047) and 57 (SL11048). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 17A and FIG. 17B. The ID₅₀ for these compounds against A549 is given in Table 2.

Example 18

Using the standard MTT protocol described above, cultured PC3 cells were exposed to serial dilutions of compounds 28 (SL11037) and 23 (SL11038). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 18A and FIG. 18B. The ID₅₀ for these compounds against PC3 is given in Table 2.

Example 19

Using the standard MTT protocol described above, cultured PC3 cells were exposed to serial dilutions of compounds 48 (SL11043) and 47 (SL11044). The percent cell

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survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 19A and FIG. 19B. The ID₅₀ for these compounds against PC3 is given in Table 2.

Example 20

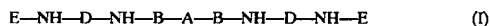
Using the standard MTT protocol described above, cultured PC3 cells were exposed to serial dilutions of compounds 58 (SL11047) and 57 (SL11048). The percent cell survival as compared to cultures exposed to vehicle alone was determined for each concentration of drug. The results are depicted in FIG. 20A and FIG. 20B. The ID₅₀ for these compounds against PC3 is given in Table 2.

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What is claimed is:

1. Compounds of Formula I:



wherein A is selected from the group consisting of cis-C₂-C₆ alkene and C₃-C₆ cycloalkyl, cycloalkenyl, and cycloaryl;

B is independently selected from the group consisting of a single bond and C₁-C₆ alkyl and alkenyl;

D is independently selected from the group consisting of C₁-C₆ alkyl and alkenyl, and C₃-C₆ cycloalkyl, cycloalkenyl, and cycloaryl;

E is methyl;

and pharmaceutically-suitable salts thereof.

2. Compounds according to claim 1, wherein A is selected from the group consisting of cyclopropyl and cyclobutyl.

3. Compounds according to claim 1, wherein B is selected from the group consisting of a single bond, methyl, and ethyl.

4. Compounds according to claim 1, wherein D is propyl.

5. Compounds according to claim 1, wherein A is selected from the group consisting of cis-2-butene, cis-cyclopropyl, trans-cyclopropyl, cis-cyclobutyl and trans-cyclobutyl; B is selected from the group consisting of a single bond and methyl; and D is propyl.

6. The compound according to claim 1, wherein A is cis-cyclopropyl, B is a single bond, and D is propyl.

7. The compound according to claim 1, wherein A is trans-cyclopropyl, B is a single bond, and D is propyl.

8. The compound according to claim 1, wherein A is cis-cyclopropyl, B is methyl, and D is propyl.

9. The compound according to claim 1, wherein A is trans-cyclopropyl, B is methyl, and D is propyl.

10. The compound according to claim 1, wherein A is trans-cyclobutyl, B is a single bond, and D is propyl.

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11. The compound according to claim 1, wherein A is cis-cyclobutyl, B is a single bond, and D is propyl.

12. The compound according to claim 1, wherein A is trans-cyclobutyl, B is methyl, and D is propyl.

13. The compound according to claim 1, wherein A is cis-cyclobutyl, B is methyl, and D is propyl. 5

14. The compound according to claim 1, wherein A is cis-2-butene, B is a single bond, and D is propyl.

15. A pharmaceutical unit dosage form for the inhibition of neoplastic cell growth comprising one or more compounds according to claim 1 in combination with a pharmaceutically-suitable carrier. 10

16. The pharmaceutical unit dosage form of claim 15, wherein the carrier is a solid carrier.

17. The pharmaceutical unit dosage form of claim 15, wherein the carrier is a liquid carrier. 15

18. A method of inhibiting growth of cancer cells comprising treating the cancer cells with an effective growth-inhibiting amount of one or more compounds according to claim 1.

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19. The method of claim 18, wherein an amount of one or more of the compounds is administered to a human cancer patient in need thereof which is effective to inhibit the growth of the cancer.

20. The method of claim 19, wherein the amount of one or more of the compounds is administered parenterally in combination with a pharmaceutically-acceptable liquid or solid carrier.

21. The method of claim 19, wherein the amount of one or more of the compounds is administered intravenously in combination with a pharmaceutically-acceptable liquid carrier.

22. The method of claim 19, wherein the amount of one or more of the compounds is administered orally in combination with a pharmaceutically-acceptable liquid or solid carrier.

* * * * *